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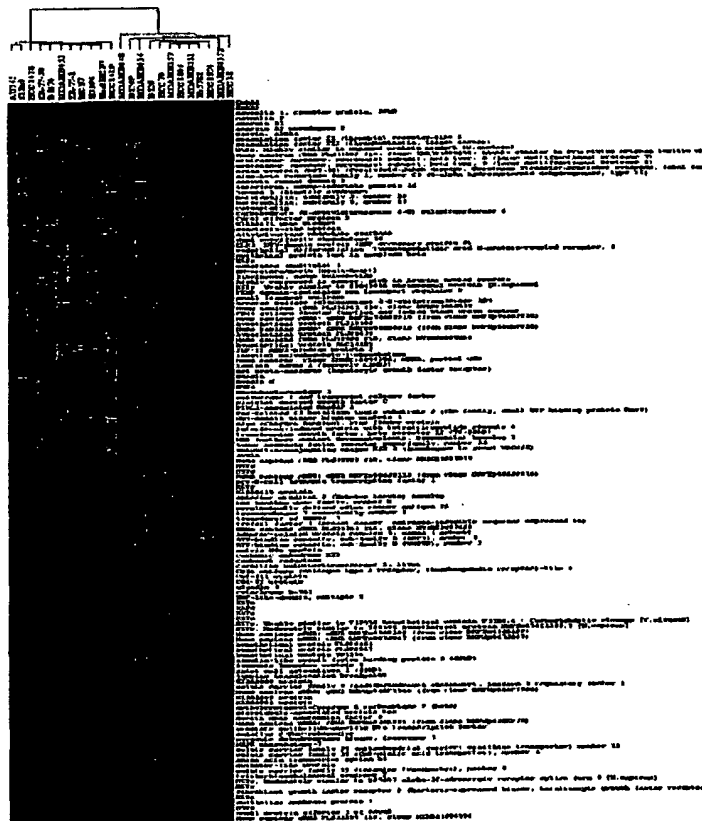
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(54) Title: IDENTIFICATION OF POLYNUCLEOTIDES FOR PREDICTING ACTIVITY OF COMPOUNDS THAT INTER-
ACT WITH AND/OR MODULATE PROTEIN TYROSINE KINASES AND/OR PROTEIN TYROSINE KINASE PATHWAYS IN
BREAST CELLS



(57) Abstract: The present invention describes polynucleotides that have been discovered to correlate to the relative intrinsic sensitivity or resistance of cells, e.g., breast cell lines, to treatment with compounds that interact with and modulate, e.g., inhibit, protein tyrosine kinases, such as, for example, members of the Src family of tyrosine kinases, e.g., Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. These polynucleotides have been shown, through a weighted voting cross validation program, to have utility in predicting the resistance and sensitivity of breast cell lines to the compounds. The expression level or phosphorylation status of some polynucleotides is regulated by treatment with a particular protein tyrosine kinase inhibitor compound, thus indicating that these polynucleotides are involved in the protein tyrosine kinase signal transduction

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pathway, e.g., Src tyrosine kinase. Such polynucleotides, whose expression levels correlate highly with drug sensitivity or resistance and which are modulated by treatment with the compounds, comprise polynucleotide predictor or marker sets useful in methods of predicting drug response, and as prognostic or diagnostic indicators in disease management, particularly in those disease areas, e.g., breast cancer, in which signaling through the protein tyrosine kinase pathway, such as the Src tyrosine kinase pathway, is involved with the disease process.

IDENTIFICATION OF POLYNUCLEOTIDES FOR PREDICTING ACTIVITY OF
COMPOUNDS THAT INTERACT WITH AND/OR MODULATE PROTEIN
TYROSINE KINASES AND/OR PROTEIN TYROSINE KINASE PATHWAYS IN
BREAST CELLS

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This application claims benefit to provisional application U.S. Serial No. 60/406,385 filed August 27, 2002, under 35 U.S.C. 119(e). The entire teachings of the referenced applications are incorporated herein by reference.

10

FIELD OF THE INVENTION

The present invention relates generally to the field of pharmacogenomics, and more specifically to new and alternative methods and procedures to determine drug sensitivity in patients, and particularly in patients with breast cancer. This invention allows the development of individualized genetic profiles which aid in treating
15 diseases and disorders based on patient response at a molecular level.

BACKGROUND OF THE INVENTION

Breast cancer is a disease with extensive histoclinical heterogeneity. Although conventional histological and clinical features have been correlated with prognosis,
20 the same apparent prognostic type of breast tumors vary widely in their responsiveness to therapy and consequent survival of the patient. New prognostic and predictive markers are needed to accurately foretell a patient's response to drugs in the clinic. Such markers would facilitate the individualization of therapy for each patient.

The problem may be solved by the identification of new parameters that can
25 better predict a patient's sensitivity to treatment or therapy. The classification of patient samples is a crucial aspect of cancer diagnosis and treatment. The association of a patient's response to drug treatment with molecular and genetic markers can open up new opportunities for drug development in non-responding patients, or distinguish a drug's indication among other treatment choices because of higher confidence in the
30 efficacy. Further, the pre-selection of patients who are likely to respond well to a medicine, drug, or combination therapy may reduce the number of patients needed in

a clinical study or accelerate the time needed to complete a clinical development program (M. Cockett et al., 2000, *Current Opinion in Biotechnology*, 11:602-609).

The major goal of pharmacogenomics research is to identify genetic markers that accurately predict a given patient's response to drugs in the clinic; such individualized genetic assessment would greatly facilitate personalized treatment. An approach of this nature is particularly needed in cancer treatment and therapy, where commonly used agents are ineffective in many patients, and side effects are frequent. The ability to predict drug sensitivity in patients is particularly challenging because drug responses reflect both the properties intrinsic to the target cells and also a host's metabolic properties. Efforts by those in the art to use genetic information to predict drug sensitivity have primarily focused on individual polynucleotides that have broad effects, such as the multidrug resistant polynucleotides, *mdr1* and *mrrp1* (P. Sonneveld, 2000, *J. Intern. Med.*, 247:521-534).

The development of microarray technologies for large scale characterization of polynucleotide expression pattern makes it possible to systematically search for multiple molecular markers and to categorize cancers into distinct subgroups that are not evident by traditional histopathological methods (J. Khan et al., 1998, *Cancer Res.*, 58:5009-5013; A.A. Alizadeh et al., 2000, *Nature*, 403:503-511; M. Bittner et al., 2000, *Nature*, 406:536-540; J. Khan et al., 2001, *Nature Medicine*, 7(6):673-679; and T.R. Golub et al., 1999, *Science*, 286:531-537; U. Alon et al., 1999, *Proc. Natl. Acad. Sci. USA*, 96:6745-6750). Such technologies and molecular tools have made it possible to monitor the expression levels of a large number of transcripts within a cell at any given time (see, e.g., Schena et al., 1995, *Science*, 270:467-470; Lockhart et al., 1996, *Nature Biotechnology*, 14:1675-1680; Blanchard et al., 1996, *Nature Biotechnology*, 14:1649; and U.S. Pat. No. 5,569,588, issued Oct. 29, 1996 to Ashby et al.).

How differential polynucleotide expression is associated with health and disease is a basis of functional genomics, which is defined as the study of all of the polynucleotides expressed by a specific cell or a group of cells and the changes in their expression pattern during development, disease, or environmental exposure. Hybridization arrays, used to study polynucleotide expression, allow polynucleotide expression analysis on a genomic scale by permitting the examination of changes in

expression of literally thousands of polynucleotides at one time. In general, for hybridization arrays, gene-specific sequences (probes) are immobilized on a solid state matrix. These sequences are then queried with labeled copies of nucleic acids from biological samples (targets). The underlying theory is that the greater the
5 expression of a gene, the greater the amount of labeled target and thus, the greater output of signal. (W.M. Freeman et al., 2000, *BioTechniques*), 29:1042-1055).

Recent studies have demonstrated that polynucleotide expression information generated by microarray analysis of human tumors can predict clinical outcome (L.J. van't Veer et al., 2002, *Nature*, 415:530-536; M. West et al., 2001, *Proc. Natl. Acad. Sci. USA*, 98:11462-11467; T. Sorlie et al., 2001, *Proc. Natl. Acad. Sci. USA*, 98:10869-10874; M. Shipp et al., 2002, *Nature Medicine*, 8(1):68-74). These findings
10 bring hope that cancer treatment will be vastly improved by better predicting the response of individual tumors to therapy.

Needed in the art are new and alternative methods and procedures to
15 determine drug sensitivity in patients and which are necessary to treat diseases and disorders, particularly cancers such as breast cancer, based on patient response at a molecular level. By using cultured cells as a model of *in vivo* effects, the present invention advantageously focuses on cell-intrinsic properties that are exposed in cell culture and involves identified polynucleotides that correlate with drug sensitivity.
20 The presently described discovery and identification of polynucleotides/marker polynucleotides (predictor polynucleotides and polynucleotide sets) in cell lines assayed *in vitro* can be used to correlate with drug responses *in vivo*, and thus can be extended to clinical situations in which the same polynucleotides are used to predict responses to drugs and/or chemotherapeutic agents by patients, with particular regard
25 to breast cancer patients.

SUMMARY OF THE INVENTION

The present invention describes the identification of marker polynucleotides whose expression levels are highly correlated with drug sensitivity in breast cell lines
30 that are either sensitive or resistant to protein tyrosine kinase inhibitor compounds. More particularly, the protein tyrosine kinases that are inhibited in accordance with the present invention include members of the Src family of tyrosine kinases, for

example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. For a review of these and other protein tyrosine kinases, see, for example, P. Blume-Jensen and T. Hunter, 2001, "Oncopolynucleotide Kinase Signaling", *Nature*, 411:355-365. Some
5 of these polynucleotides are also modulated by the tyrosine kinase inhibitor compounds, in particular, src tyrosine kinase inhibitor compounds, which indicates their involvement in the protein tyrosine kinase signaling pathway. These polynucleotides or "markers" show utility in predicting a host's response to a drug and/or drug treatment. Similar expression pattern of these polynucleotides to breast
10 cell lines is also seen in primary breast tumors which indicates co-regulation of these marker polynucleotides.

It is an aspect of this invention to provide a cell culture model to identify polynucleotides whose expression levels correlate with drug sensitivity of cells associated with a disease state, or with a host having a disease. In accordance with the
15 present invention, oligonucleotide microarrays were utilized to measure the expression levels of a large number of polynucleotides in a panel of untreated cell lines, particularly breast cell lines, for which drug sensitivity to a protein tyrosine kinase inhibitor compound was determined. The determination of the polynucleotide expression profiles in the untreated cells allowed a prediction of chemosensitivity and
20 the identification of marker polynucleotides whose expression levels highly correlated with sensitivity to drugs or compounds that modulate, preferably inhibit, protein tyrosine kinase or the pathway in which the protein tyrosine kinase, e.g., src tyrosine kinase, is involved. The marker polynucleotides are thus able to be utilized as one or more predictors to foresee a patient's response to drugs or drug treatments that
25 directly or indirectly affect protein tyrosine kinase activity.

It is another aspect of the present invention to provide a method of determining or predicting if an individual requiring drug or chemotherapeutic treatment or therapy for a disease state, or a cancer or tumor of a particular type, e.g., a breast cancer or breast tumor, will successfully respond or will not respond to the
30 drug or chemotherapeutic treatment or therapy prior to the administration of such treatment or chemotherapy. Preferably, the treatment or therapy involves a protein tyrosine kinase modulating agent, e.g., an inhibitor of the protein tyrosine kinase

activity. The protein tyrosine kinases whose activities can be inhibited by inhibitor compounds according to this invention include, for example, members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Also in accordance with the present invention, cells from a patient tissue sample, e.g., a breast tumor or cancer biopsy, are assayed to determine their polynucleotide expression pattern prior to treatment with a protein tyrosine kinase modulating compound or drug, preferably a src tyrosine kinase inhibitor. The resulting polynucleotide expression profile of the test cells before exposure to the compound or drug is compared with the polynucleotide expression pattern of the predictor set of polynucleotides that have been described and shown herein (Table 2). In addition, in such a method, the polynucleotide expression pattern of subsets of predictor polynucleotides, i.e., the sets of 15 and 7 polynucleotides as set forth in Tables 4-5, respectively, can also be used. These polynucleotides are derived from the control panel of the untreated cells that have been determined to be either resistant or sensitive to the drug or compound, i.e., FIG. 1 and Table 1.

Success or failure of treatment with a drug can be determined based on the polynucleotide expression pattern of cells from the test tissue (test cells), e.g., a tumor or cancer biopsy, as being relatively similar to or different from the polynucleotide expression pattern of the predictor set of polynucleotides. Thus, if the test cells show a polynucleotide expression profile which corresponds to that of the predictor set of polynucleotides in the control panel of cells which are sensitive to the drug or compound, it is highly likely or predicted that the individual's cancer or tumor will respond favorably to treatment with the drug or compound. By contrast, if the test cells show a polynucleotide expression pattern corresponding to that of the predictor set of polynucleotides of the control panel of cells which are resistant to the drug or compound, it is highly likely or predicted that the individual's cancer or tumor will not respond to treatment with the drug or compound.

It is a further aspect of this invention to provide screening assays for determining if a cancer patient will be susceptible or resistant to treatment with a drug or compound, particularly, a drug or compound directly or indirectly involved in a protein tyrosine kinase activity or a protein tyrosine kinase pathway. Such protein

tyrosine kinases include, without limitation, members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors.

5 In a more particular aspect, the present invention provides screening assays for determining if a cancer patient will be susceptible or resistant to treatment with a drug or compound, particularly, a drug or compound directly or indirectly involved in src tyrosine kinase activity or the src tyrosine kinase pathway.

It is another aspect of the present invention to provide a method of monitoring the treatment of a patient having a disease treatable by a compound or agent that
10 modulates a protein tyrosine kinase, including members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. This can be accomplished by comparing the resistance or sensitivity polynucleotide expression profile of cells from a patient tissue sample, e.g., a tumor or cancer biopsy, e.g., a breast cancer or tumor sample, prior to treatment with a drug or compound that
15 inhibits the protein tyrosine kinase activity and again following treatment with the drug or compound. The isolated test cells from the patient's tissue sample are assayed to determine their polynucleotide expression pattern before and after exposure to a compound or drug, such as, e.g., a src tyrosine kinase inhibitor. The resulting
20 polynucleotide expression profile of the test cells before and after treatment is compared with the polynucleotide expression pattern of the predictor set and subsets of polynucleotides that have been described and shown herein to be highly expressed in the control panel of cells that are either resistant or sensitive to the drug or compound. Thus, if a patient's response becomes one that is sensitive to treatment by
25 a protein tyrosine kinase inhibitor compound, based on a correlation of the expression profile of the predictor polynucleotides, the patient's treatment prognosis can be qualified as favorable and treatment can continue. Also, if after treatment with a drug or compound, the test cells do not show a change in their polynucleotide expression profile that corresponds to the control panel of cells that are sensitive to the drug or
30 compound, this can serve as an indicator that the current treatment should be modified, changed, or even discontinued. Such a monitoring process can indicate

success or failure of a patient's treatment with a drug or compound, and the monitoring processes can be repeated as necessary or desired.

It is a further aspect of the present invention to provide predictor polynucleotides and predictor sets of polynucleotides having both diagnostic and prognostic value in disease areas in which signaling through a protein tyrosine kinase or a protein tyrosine kinase pathway is of importance, e.g., in cancers and tumors, in immunological disorders, conditions or dysfunctions, or in disease states in which cell signaling and/or proliferation controls are abnormal or aberrant. Such protein tyrosine kinases whose direct or indirect modulation can be associated with a disease state or condition, include members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. In accordance with this invention, the use of predictor polynucleotides, or a predictor polynucleotide set or subset (such as the predictor polynucleotides of Table 2, and the predictor polynucleotide subsets of Tables 4-5) is to forecast or foretell an outcome prior to having any knowledge about a biological system, or a cellular response.

It is yet another aspect of the present invention to assemble polynucleotides, such as those listed in Table 2, or the subset of polynucleotides as listed in Tables 4-5, that highly correlate with resistance or sensitivity to protein tyrosine kinase inhibitor drugs or compounds, into predictor polynucleotide sets, so as to predict, or reasonably foretell the effect of either the protein tyrosine inhibitor compounds, or compounds that affect the protein tyrosine kinase signaling pathway(s) in different biological systems, or for cellular responses. The predictor polynucleotide sets can be used in *in vitro* assays of drug response by test cells to predict *in vivo* outcome. In accordance with this invention, the various predictor polynucleotide sets described herein, or the combination of these predictor sets with other polynucleotides or other co-variants of these polynucleotides, can be used, for example, to predict how patients with cancer or a tumor might respond to therapeutic intervention with compounds that modulate protein tyrosine kinases, or modulate signaling through an entire protein tyrosine kinase regulatory pathway. The predictor sets of polynucleotides, or co-variants of these polynucleotides, can be used to predict how patients with a cancer or tumor respond to therapy employing compounds that modulate a tyrosine kinase, or the

activity of a tyrosine kinase, such as protein tyrosine kinase members of the Src family, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors.

Another object of the present invention is to provide one or more specialized
5 microarrays, e.g., oligonucleotide microarrays or cDNA microarrays, comprising those polynucleotides or combinations thereof, as described herein, showing expression profiles that correlate with either sensitivity or resistance to protein tyrosine kinase inhibitor compounds. Such microarrays can be employed in *in vitro* assays for assessing the expression level of the polynucleotides on the microarrays in
10 the test cells from tumor biopsies, for example, and determining whether these test cells will be likely to be resistant or sensitive to the protein tyrosine kinase inhibitor compound(s). For example, a specialized microarray can be prepared using some or all of the polynucleotides, polynucleotide subsets, or combinations thereof, as described herein and shown in Tables 2, 4 and 5. Cells from a tissue or organ biopsy
15 can be isolated and exposed to one or more inhibitor compounds. Following application of nucleic acids isolated from both untreated and treated cells to one or more of the specialized microarrays, the pattern of polynucleotide expression of the tested cells can be determined and compared with that of the predictor polynucleotide pattern from the control panel of cells used to create the predictor polynucleotide set
20 on the microarray. Based upon the polynucleotide expression pattern results from the cells undergoing testing, it can be determined if the cells show a resistant or a sensitive profile of polynucleotide expression. Whether or not the tested cells from a tissue or organ biopsy will respond to a protein tyrosine kinase inhibitor compound, and the course of treatment or therapy, can then be determined or evaluated based on
25 the information gleaned from the results of the specialized microarray analysis.

It is a further aspect of the present invention to provide a kit for determining or predicting drug susceptibility or resistance by a patient having a disease, with particular regard to a cancer or tumor, namely, a breast cancer or tumor. Such kits are useful in a clinical setting for testing a patient's biopsied tumor or cancer sample, for
30 example, to determine or predict if the patient's tumor or cancer will be resistant or sensitive to a given treatment or therapy with a drug, compound, chemotherapy agent, or biological agent that is directly or indirectly involved with modification, preferably,

inhibition, of the activity of a protein tyrosine kinase or a cell signaling pathway involving protein tyrosine kinase activity. Provided in the kit are one or more microarrays, e.g., oligonucleotide microarrays or cDNA microarrays, comprising those polynucleotides that correlate with resistance and sensitivity to protein tyrosine

5 kinase modulators, particularly, inhibitors of members of the Src family of protein tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as inhibitors of the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases; and, in suitable containers, the modulator agents/compounds for use in testing cells from patient tissue specimens or patient samples; and instructions for use.

10 In addition, kits contemplated by the present invention can include reagents or materials for the monitoring of the expression of the predictor or marker polynucleotides of the invention at the level of mRNA or protein, using other techniques and systems practiced in the art, e.g., RT-PCR assays, which employ primers designed on the basis of one or more of the predictor polynucleotides

15 described herein, immunoassays, such as enzyme linked immunosorbent assays (ELISAs), immunoblotting, e.g., Western blots, or *in situ* hybridization, and the like, as further described herein. The kits according to the present invention can also comprise predictor polynucleotides as set forth in Table 2, and/or one or more of the predictor polynucleotide subsets as presented in Tables 4-5 herein.

20 Another aspect of the present invention is to provide one or more polynucleotides among those of the predictor polynucleotides identified herein that can serve as targets for the development of drug therapies for disease treatment. Such targets can be particularly applicable to treatment of breast disease, such as breast cancers or tumors. Because these predictor polynucleotides are differentially

25 expressed in sensitive and resistant cells, their expression pattern is correlated with the relative intrinsic sensitivity of cells to treatment with compounds that interact with and/or inhibit protein tyrosine kinases, including members of the Src family of protein tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases.

30 Accordingly, the polynucleotides highly expressed in resistant cells can serve as targets for the development of new drug therapies for those tumors which are resistant to protein tyrosine kinase inhibitor compounds.

Yet another object of the present invention is to provide antibodies, either polyclonal or monoclonal, directed against one or more of the protein tyrosine kinase biomarker polypeptides, or peptides thereof, encoded by the predictor polynucleotides. Such antibodies can be used in a variety of ways, for example, to
5 purify, detect, and target the protein tyrosine kinase biomarker polypeptides of the present invention, including both *in vitro* and *in vivo* diagnostic, detection, screening, and/or therapeutic methods, and the like. Included among the protein tyrosine kinase biomarker polypeptides of this invention are members of the Src family of protein tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as
10 the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases.

Yet another object of the present invention is to provide antisense reagents, including siRNA, RNAi, and ribozyme reagents, directed against one or more of the protein tyrosine kinase biomarker polypeptides, or peptides thereof, encoded by the predictor polynucleotides. Such antisense reagents can be used in a variety of ways,
15 for example, to detect, to target, and inhibit the expression of the protein tyrosine kinase biomarker polypeptides of the present invention, including both *in vitro* and *in vivo* diagnostic, detection, screening, and/or therapeutic methods, and the like. Included among the protein tyrosine kinase biomarker polypeptides of this invention are members of the Src family of protein tyrosine kinases, for example, Src, Fgr, Fyn,
20 Yes, Blk, Hck, Lck and Lyn, as well as the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases.

The invention also relates to an antisense compound 8 to 30 nucleotides in length that specifically hybridizes to a nucleic acid molecule encoding the human protein tyrosine kinase biomarker polypeptides of the present invention, wherein said
25 antisense compound inhibits the expression of the human protein tyrosine kinase biomarker polypeptides.

The invention further relates to a method of inhibiting the expression of the human protein tyrosine kinase biomarker polypeptides of the present invention in human cells or tissues comprising contacting said cells or tissues *in vitro*, or *in vivo*,
30 with an antisense compound of the present invention so that expression of the protein tyrosine kinase biomarker polypeptides is inhibited.

The present invention is also directed to a method of identifying a compound that modulates the biological activity of protein tyrosine kinase biomarker polypeptides, comprising the steps of, (a) combining a candidate modulator compound with protein tyrosine kinase biomarker polypeptides in the presence of an
5 antisense molecule that antagonizes the activity of the protein tyrosine kinase biomarker polypeptides selected from the group consisting of SEQ ID NO:534 thru 557, and (b) identifying candidate compounds that reverse the antagonizing effect of the peptide.

The present invention is also directed to a method of identifying a compound
10 that modulates the biological activity of protein tyrosine kinase biomarker polypeptides, comprising the steps of, (a) combining a candidate modulator compound with protein tyrosine kinase biomarker polypeptides in the presence of a small molecule that antagonizes the activity of the protein tyrosine kinase biomarker polypeptides selected from the group consisting of SEQ ID NO:534 thru 557, and (b)
15 identifying candidate compounds that reverse the antagonizing effect of the peptide.

The present invention is also directed to a method of identifying a compound that modulates the biological activity of protein tyrosine kinase biomarker polypeptides, comprising the steps of, (a) combining a candidate modulator compound with protein tyrosine kinase biomarker polypeptides in the presence of a
20 small molecule that agonizes the activity of the protein tyrosine kinase biomarker polypeptides selected from the group consisting of SEQ ID NO:534 thru 557, and (b) identifying candidate compounds that reverse the agonizing effect of the peptide.

Further aspects, features, and advantages of the present invention will be better appreciated upon a reading of the detailed description of the invention when
25 considered in connection with the accompanying figures or drawings.

DESCRIPTION OF THE FIGURES

The file of this patent contains at least one Figure executed in color. Copies of this patent with color Figure(s) will be provided by the Patent and Trademark Office
30 upon request and payment of the necessary fee.

FIG. 1 illustrates a polynucleotide expression pattern according to the present invention. The 137 polynucleotides that highly correlated with a resistance/sensitivity

phenotype classification of the 23 breast cell lines for the protein tyrosine kinase inhibitor compound BMS-A are shown. Each row corresponds to a polynucleotide, with the columns corresponding to expression level in the different cell lines. Expression levels for each polynucleotide were normalized across all 23 breast cell lines such that the median is 0 and the standard derivation is 1. The expression levels greater than the median are shaded in red, and those below the mean are shaded in green. The individual polynucleotides encoding the protein tyrosine kinase biomarkers of the invention are indicated at the right (details of the biomarkers are also shown in the Table 2). The cell lines labeled in red are classified as resistant, and those labeled in blue are classified as sensitive to BMS-A according to their IC₅₀.

FIG. 2 The examples of polynucleotides whose expression levels are not only correlated with the sensitivity or resistance of breast cell lines to treatment with a protein tyrosine kinase inhibitor compound (e.g., BMS-A), but also differentially down regulated by treatment with the compound. Eleven breast cell lines (5 sensitive and 6 resistant cell lines as indicated in bold in the Table 1) were used in a drug treatment study. Cells were treated with or without the BMS-A compound (0.4 μ M) in 0.1% DMSO for 24 hours. Expression profiling was performed, the polynucleotide expression of a cell line treated with drug was compared pair-wisely to the polynucleotide expression of the same cell line without drug treatment. Five sensitive cell lines without drug treatment are indicated with lightly shaded bars ("A" side of graph); five sensitive cell lines with drug treatment are indicated in darkly shaded bars ("A" side of graph); six resistant cell lines without drug treatment are indicated in darkly shaded bars ("B" side of graph); six resistant cell lines with drug treatment are indicated in lightly shaded bars ("B" side of graph).

FIG. 3 The examples of polynucleotide whose expression is down regulated by BMS-A compound treatment in a dose and time dependent manner in a prostate cell line PC3. Cells are treated without or with 0.025 μ M, 0.1 μ M and 0.4 μ M of the BMS-A compound for 4 hours or 24 hours. The relative polynucleotide expression level of treated cells is compared to the corresponding untreated control which is set to 1. Drug concentrations and time of treatment are indicated.

FIG. 4 Immunoblot analysis of EphA2 protein level and tyrosine phosphorylation status in nine breast tumor cell lines. Cells were treated with 0.1 μ M

BMS-A for 1 hour. Cell lysates were immuno-precipitated with EphA2 antibody and blotted with EphA2 antibody (to assess EphA2 protein level) or anti-phosphotyrosine antibody (to assess EphA2 tyrosine phosphorylation status). Cell lines with or without drug treatment are indicated. The results indicate that EphA2 protein level does not
5 change upon one hour drug treatment, but the phosphorylation of tyrosine residues is dramatically decreased with the drug treatment.

FIG. 5 shows the error rates of different predictor sets comprising the marker polynucleotides with differential selection and combination for the BMS-A protein tyrosine kinase inhibitor compound in the leave-one-out cross validation tests. The
10 Genecluster software was used to select polynucleotides and predict classifications using a "weighted-voting leave-one-out cross-validation algorithm", as described herein. A different number of polynucleotides was selected in the predictor set from (i) the 137 polynucleotides, or (ii) the 40 polynucleotides modulated by BMS-A treatment as shown in Table 2, for predicting resistant and sensitive classes to BMS-A
15 in the breast cell lines. FIG. 5 demonstrates that a different selection and different combination of polynucleotides in a predictor set achieve different error rates in the leave-one-out cross validation. When the predictor sets were selected from 137 polynucleotides as shown in Table 2, the lowest error rate of 6.3% was achieved in the leave-one-out cross validation with 15 markers. Another predictor set comprised of 7
20 polynucleotides selected from the 40 polynucleotides that were modulated by the drug treatment achieved an error rate of 3.1%. These results indicate that polynucleotides which are not only correlated with drug sensitivity, but also modulated by the drug, can provide a better and more accurate prediction in a predictor set.

FIG. 6 shows the error rate comparison for predicting the sensitivity
25 classification of compound BMS-A in the breast cell lines and random permutation tests in leave-one-out cross validation. When a predictor set contained either 7 or 15 polynucleotides selected from different polynucleotide groups, the error rate of the leave-one-out cross validation tests for predicting sensitivity of BMS-A in the 23 breast cell lines was 3.1% and 6.3% respectively. In contrast, the real error rates
30 ranged from 30% to 83% when the same number of polynucleotides in a predictor set was used in 20 cases in which classification for the breast cell lines was randomly assigned. This result demonstrates that the error rate value for predicting sensitivity

of BMS-A in the 23 breast cell lines is significantly lower than the error rate for predicting sensitivity for the 23 breast cell lines when their classification is randomly assigned in 20 cases.

FIG. 7 The expression pattern of the 137 marker polynucleotides in 134 primary breast tumors. These 137 polynucleotides are highly correlated with a resistance/sensitivity phenotype classification of the 23 breast cell lines for the protein tyrosine kinase inhibitor compound BMS-A according to the present invention (as shown in FIG.1). Each row corresponds to a gene, with the columns corresponding to expression level in the different breast tumor samples. Expression levels for each polynucleotide were normalized across all 134 breast tumor samples such that the median is 0 and the standard derivation is 1. The expression levels greater than the median are shaded in red, and those below the mean are shaded in green. The order of individual polynucleotides encoding the protein tyrosine kinase biomarkers of the invention are the same as indicated in FIG.1. The expression pattern clearly shows that a group of primary breast tumors (as indicated by the arrow) highly expressed sensitive markers of protein tyrosine kinase inhibitor compound of the invention. By contrast, another different group highly expressed resistant markers.

DESCRIPTION OF THE TABLES

Table 1 presents the resistance/sensitivity phenotype classification of the 23 breast cell lines for the protein tyrosine kinase inhibitor compound BMS-A based on IC_{50} results. The IC_{50} for each cell line was assessed in by MTS assays as described in Example 1 (Methods). The mean IC_{50} values along with standard deviations (SD) were calculated from 2 to 5 individual determinations for each cell line as shown. The IC_{50} unit is μM . The mean IC_{50} for each cell line was log-transformed to $\log_{10}(IC_{50})$ and the mean $\log_{10}(IC_{50})$ across the 23 breast cell lines for BMS-A was calculated and used to normalize the IC_{50} data for each cell line. The cell lines with a $\log_{10}(IC_{50})$ below the mean $\log_{10}(IC_{50})$ were defined as sensitive to the compound, while those having a $\log_{10}(IC_{50})$ above the mean $\log_{10}(IC_{50})$ were considered to be resistant. The cell lines presented in bold were used in the drug induction study as described herein.

TABLE 1

#	Cell Lines	mean IC ₅₀ (μM) to BMS-A	SD	Log(IC ₅₀)	Normalized Log(IC ₅₀)	Classification
1	MDA-MB-157	0.0055	0.0035	-2.25924	-2.25405	Sensitive
2	MDA-MB-231	0.0095	0.0058	-2.02422	-2.03843	Sensitive
3	HCC1954	0.0242	0.0172	-1.61621	-1.66411	Sensitive
4	HCC70	0.0337	0.0160	-1.47214	-1.53193	Sensitive
5	BT-20	0.1652	0.1036	-0.78195	-0.89871	Sensitive
6	HCC1806	0.2194	0.1508	-0.65884	-0.78576	Sensitive
7	HS578T	0.6472	0.5885	-0.18898	-0.35469	Sensitive
8	HCC1419	2.5093	0.2280	0.399548	0.18525	Resistant
9	SK-BR-3	2.7534	0.8410	0.439867	0.22224	Resistant
10	AU-565	5.2399	3.2627	0.719322	0.47863	Resistant
11	HCC38	6.6327	3.1673	0.821688	0.57254	Resistant
12	BT-474	6.7375	4.1515	0.828502	0.57880	Resistant
13	MDA-MB-468	7.1258	4.0960	0.852833	0.60112	Resistant
14	HCC1428	7.2926	4.1436	0.862881	0.61034	Resistant
15	MDA-MB-435S	7.7800	2.3643	0.89098	0.63612	Resistant
16	H3396	8.1950	3.2549	0.91355	0.65682	Resistant
17	BT-549	9.0576	1.1419	0.957014	0.69670	Resistant
18	ZR-75-30	9.2632	0.5827	0.966762	0.70564	Resistant
19	MCF7	>9.5238	1.95E-07	0.978811	0.71670	Resistant
20	MCF7/Her2	>9.5238	1.8E-07	0.978811	0.71670	Resistant
21	MDA-MB-436	>9.5238	1.51E-07	0.978811	0.71670	Resistant
22	ZR-75-1	>9.5238	1.8E-07	0.978811	0.71670	Resistant
23	MDA-MB-453	>9.5238		0.978811	0.71670	Resistant
	Mean IC ₅₀ across all 23 cell lines	5.2744		0.197626		
	SD	3.9565		1.08998		

Table 2 shows a polynucleotide list derived from three analysis algorithms that demonstrated a high correlation between expression pattern and resistance/sensitivity classification to BMS-A. The polynucleotide number, relative expression pattern, i.e., sensitive or resistant, Genbank Accession number, polynucleotide description, Unigene cluster number, SEQ ID NO: for the nucleic acid sequence of the gene, SEQ ID NO: for the amino acid sequence coded for by the polynucleotide (if available) and PID (protein ID), are presented in Table 2. For each gene, the DNA and encoded amino acid sequence represented by SEQ ID NOs. in Table 2 are set forth in the Sequence Listing.

TABLE 2
Markers highly correlated to BMS-A in expression pattern and resistance/sensitivity classification

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
1	Sensitive cells	NM_004431	yes	EphA2	Hs.171596	1	138	NP_004422
2	Sensitive cells	AF025304		EphB2	Hs.125124	2	139	AAB94602
3	Sensitive cells	AU147399	yes	caveolin 1, caveolae protein, 22kD	Hs.74034	3	140	NP_001744
4	Sensitive cells	NM_001233	yes	caveolin 2	Hs.139851	4	141	NP_001224
5	Sensitive cells	NM_000700	yes	annexin A1	Hs.78225	5	142	NP_000691
6	Sensitive cells	NM_004039		annexin A2	Hs.406239	6	143	NP_004030
7	Sensitive cells	BG107577		parvin, alpha	Hs.44077	7	144	Q9NVD7
8	Sensitive cells	BE965369	yes	coagulation factor II (thrombin) receptor-like 1	Hs.154299	8	145	XP_003671
9	Sensitive cells	NM_001993	yes	coagulation factor III (thromboplastin, tissue factor)	Hs.62192	9	146	NP_001984
10	Sensitive cells	BF792126		Homo sapiens, clone IMAGE:4344858, mRNA	Hs.432974	10	147	P1_453619
11	Sensitive cells	BE856341		layilin	Hs.133015	11	148	Q96NF3
12	Sensitive cells	U17496		proteasome (prosome, macropain) subunit, beta type, 8 (large multifunctional protease 7)	Hs.180062	12	149	P28062
13	Sensitive cells	NM_002800		proteasome (prosome, macropain) subunit, beta type, 9 (large multifunctional protease 2)	Hs.381081	13	150	NP_002791
14	Sensitive cells	NM_000311		prion protein (p27-30) (Creutzfeld-Jakob disease, Gerstmann-Strausler-Scheinker syndrome, fatal familial insomnia)	Hs.74621	14	151	P04156
15	Sensitive cells	NM_003739	yes	aldo-keto reductase family 1, member C3 (3-alpha hydroxysteroid dehydrogenase, type II)	Hs.78183	15	152	NP_003730
16	Sensitive cells	NM_020639		ankyrin repeat domain 3	Hs.55565	16	153	NP_065690
17	Sensitive cells	AF208043	yes	interferon, gamma-inducible protein 16	Hs.155530	17	154	Q16666
18	Sensitive cells	AF003837	yes	jagged 1 (Alagille syndrome)	Hs.91143	18	155	P78504
19	Sensitive cells	BC002832	yes	butyrophilin, subfamily 3, member A2	Hs.87497	19	156	AAF76140
20	Sensitive cells	NM_006994	yes	butyrophilin, subfamily 3, member A3	Hs.167741	20	157	NP_008925
21	Sensitive cells	AF327443		calpastatin	Hs.359682	21	158	XP_051211

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
22	Sensitive cells	NM_021615	yes	carbohydrate (N-acetylglucosamine 6-O) sulfotransferase 6	Hs.157439	22	159	NP_067628
23	Sensitive cells	AF104857	yes	CDC42 effector protein (Rho GTPase binding) 3	Hs.260024	23	160	NP_006440
24	Sensitive cells	AL136896		suppressor of cytokine signaling 5	Hs.169836	24	161	O75159
25	Sensitive cells	AL565621	yes	coactosin-like protein	Hs.289092	25	162	AAH16702
26	Sensitive cells	BF111719		alkylglycerone phosphate synthase	Hs.22580	26	163	O00116
27	Sensitive cells	N36770		dual specificity phosphatase 10	Hs.177534	27	164	NP_009138
28	Sensitive cells	AW575374	yes	ELK3, ETS-domain protein (SRF accessory protein 2)	Hs.288555	28	165	NP_005221
29	Sensitive cells	AW269335	yes	endothelial differentiation, lysophosphatidic acid G-protein-coupled receptor, 2	Hs.75794	29	166	NP_001392
30	Sensitive cells	BC001247	yes	epithelial protein lost in neoplasm beta	Hs.10706	30	167	Q9UHB6
31	Sensitive cells	BE669858		hypothetical protein FLJ39885	Hs.319825	31	168	NP_689916
32	Sensitive cells	NM_000127		exostoses (multiple) 1	Hs.184161	32	169	NP_000118
33	Sensitive cells	NM_002589		BH-protocadherin (brain-heart)	Hs.34073	33	170	O60245
34	Sensitive cells	AI133452		fibrinogen, gamma polypeptide	Hs.75431	34	171	AAH21674
35	Sensitive cells	NM_006101		highly expressed in cancer, rich in leucine heptad repeats	Hs.58169	35	172	NP_006092
36	Sensitive cells	AL135264		ESTs, Moderately similar to hypothetical protein FLJ20489	Hs.406100	36		
37	Sensitive cells	NM_014164		FXYD domain-containing ion transport regulator 5	Hs.333418	37	173	NP_054883
38	Sensitive cells	BC003502		small fragment nuclease	Hs.7527	38	174	Q9Y3B8
39	Sensitive cells	AA780067		heparan sulfate (glucosamine) 3-O-sulfotransferase 3B1	Hs.159572	39	175	Q9Y662
40	Sensitive cells	AA702248	yes	Homo sapiens cDNA FLJ14241 fis, clone OVARC1000533	Hs.183765	40		
41	Sensitive cells	BC004372		CD44 antigen (homing function and Indian blood group system)	Hs.169610	41	176	Q9UJ36
42	Sensitive cells	BF688144		Homo sapiens mRNA; cDNA DKFZp762O2215 (from clone DKFZp762O2215)	Hs.331666	42		
43	Sensitive cells	NM_018067		hypothetical protein FLJ10350	Hs.177596	43	177	NP_060537
44	Sensitive cells	BG111761		guanine nucleotide binding protein (G protein), gamma 12	Hs.8107	44	178	Q9UB16
45	Sensitive cells	NM_017821		nucleoredoxin	Hs.374534	45	179	NP_060291

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
46	Sensitive cells	AA722799		endothelial and smooth muscle cell-derived neuropilin-like protein	Hs.173374	46	180	Q96PD2
47	Sensitive cells	BC006436		hypothetical protein MGC13105	Hs.22744	47	181	AAH06436
48	Sensitive cells	NM_006548	yes	IGF-II mRNA-binding protein 2	Hs.30299	48	182	NP_006539
49	Sensitive cells	NM_002194		inositol polyphosphate-1-phosphatase	Hs.32309	49	183	NP_002185
50	Sensitive cells	BC251556		KIAA1949 protein	Hs.101150	50	184	BAH85535
51	Sensitive cells	J03202		laminin, gamma 1 (formerly LAMB2)	Hs.432855	51	185	NP_002284
52	Sensitive cells	NM_000245	yes	met proto-oncogene (hepatocyte growth factor receptor)	Hs.316752	52	186	NP_000236
53	Sensitive cells	NM_002444		moesin	Hs.170328	53	187	NP_002435
54	Sensitive cells	NM_012334	yes	myosin X	Hs.61638	54	188	NP_036466
55	Sensitive cells	AI769569		ESTs	Hs.112472	55		
56	Sensitive cells	NM_002633	yes	phosphoglucomutase 1	Hs.1869	56	189	NP_002624
57	Sensitive cells	BC004295	yes	polymerase I and transcript release factor	Hs.29759	57	190	O00535
58	Sensitive cells	NM_016205		platelet derived growth factor C	Hs.43080	58	191	Q9UL22
59	Sensitive cells	NM_004815	yes	PTPL1-associated RhoGAP 1	Hs.70983	59	192	NP_004806
60	Sensitive cells	NM_002872		ras-related C3 botulinum toxin substrate 2 (rho family, small GTP binding protein Rac2)	Hs.301175	60	193	NP_002863
61	Sensitive cells	AF329267		SH3-domain kinase binding protein 1	Hs.153260	61	194	XP_039010
62	Sensitive cells	AI572079		snail homolog 2 (Drosophila)	Hs.93005	62	195	AAH14890
63	Sensitive cells	NM_001549		interferon-induced protein with tetratricopeptide repeats 4	Hs.181874	63	196	O14879
64	Sensitive cells	D50683		transforming growth factor, beta receptor II (70-80kD)	Hs.82028	64	197	NP_003233
65	Sensitive cells	NM_005902		MAD (mothers against decapentaplegic, Drosophila) homolog 3	Hs.288261	65	198	Q92940
66	Sensitive cells	NM_014452		tumor necrosis factor receptor superfamily, member 21	Hs.159651	66	199	NP_05267
67	Sensitive cells	AB017644		ubiquitin-conjugating enzyme E2E 3 (homologous to yeast UBC4/5)	Hs.4890	67	200	XP_096160
68	Sensitive cells	BC002323		zyxin	Hs.75873	68	201	Q15942
69	Resistant cells	AL157452		Homo sapiens mRNA; cDNA DKFZp761C1712 (from clone DKFZp761C1712)	Hs.4774	69		

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
70	Resistant cells	BF752277		hypothetical protein FLJ20151	Hs.279916	70	202	Q9NXM9
71	Resistant cells	BF512299		ESTs	Hs.438672	71		
72	Resistant cells	AL049381	yes	Homo sapiens mRNA; cDNA DKFZp586j2118 (from clone DKFZp586j2118)	Hs.21851	72		
73	Resistant cells	NM_002585	yes	pre-B-cell leukemia transcription factor 1	Hs.155691	73	203	NP_002576
74	Resistant cells	T68445		anaphase-promoting complex subunit 7	Hs.52763	74	204	Q96AC4
75	Resistant cells	BF308645		PRx1 KIAA1415 protein	Hs.109315	75	205	Q8TCU6
76	Resistant cells	AF088867	yes	anterior gradient 2 (Xenopus laevis) homolog	Hs.413945	76	206	AF088867_1
77	Resistant cells	NM_004040	yes	Human HepG2 3' region cDNA, clone hmd1f06.	Hs.204354	77	207	NP_004031
78	Resistant cells	AF151810	yes	serologically defined colon cancer antigen 28	Hs.84700	78	208	Q9Y365
79	Resistant cells	NM_004252		transmembrane 7 superfamily member 2	Hs.31130	79	209	NP_004243
80	Resistant cells	NM_005749	yes	transducer of ERBB2, 1	Hs.178137	80	210	NP_005740
81	Resistant cells	NM_003225	yes	trefoil factor 1 (breast cancer, estrogen-inducible sequence expressed in)	Hs.350470	81	211	NP_003216
82	Resistant cells	AA181060	yes	Homo sapiens cDNA FLJ31753 fis, clone NT2RI2007468	Hs.349283	82		
83	Resistant cells	AL050025		adaptor-related protein complex 1, gamma 1 subunit	Hs.5344	83	212	CAB43244
84	Resistant cells	NM_001089		ATP-binding cassette, sub-family A (ABC1), member 3	Hs.26630	84	213	NP_001080
85	Resistant cells	NM_004915		ATP-binding cassette, sub-family G (WHITE), member 1	Hs.10237	85	214	NP_004906
86	Resistant cells	AL523275		CALM1 calmodulin 1 (phosphorylase kinase, delta)	Hs.374441	86	215	AAH00454
87	Resistant cells	NM_001218	yes	carbonic anhydrase XII	Hs.5338	87	216	NP_001209
88	Resistant cells	NM_016286		dicarbonyl/L-xylulose reductase	Hs.9857	88	217	NP_057370
89	Resistant cells	BC000185		carnitine palmitoyltransferase I, liver	Hs.259785	89	218	AAH00185
90	Resistant cells	NM_005505		scavenger receptor class B, member 1	Hs.180616	90	219	NP_005496
91	Resistant cells	NM_016048		CGI-111 protein	Hs.11085	91	220	NP_057132
92	Resistant cells	BC000195		CGI-81 protein	Hs.279583	92	221	NP_057109
93	Resistant cells	NM_001306		claudin 3	Hs.25640	93	222	NP_001297
94	Resistant cells	BC000021		cytochrome b-561	Hs.355264	94	223	NP_001906
95	Resistant cells	W68084		EGF-like-domain, multiple 5	Hs.5599	95	224	Q9HIU4
96	Resistant cells	AA825563	yes	ESTs	Hs.445708	96		

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
97	Resistant cells	BE887449		Homo sapiens cDNA FLJ34170 fis, clone FCBBF3015396.	Hs.32112	97		
98	Resistant cells	AI123815	yes	hypothetical protein FLJ21963	Hs.13222	98	225	Q9H6R3
99	Resistant cells	AI308862		RAB21, member RAS oncogene family	Hs.184627	99	226	Q9UL25
100	Resistant cells	AW006352		EST	Hs.159643	100		
101	Resistant cells	AL554277		chromosome 17 open reading frame 28	Hs.11067	101	227	Q9NT34
102	Resistant cells	BG289001		hypothetical protein LOC253782	Hs.387400	102		
103	Resistant cells	AI935915		hypothetical protein LOC112868	Hs.97837	103	228	XP_053402
104	Resistant cells	NM_017689	yes	hypothetical protein FLJ20151	Hs.279916	104	229	NP_060159
105	Resistant cells	NM_017966		hypothetical protein FLJ20847	Hs.13479	105	230	NP_060436
106	Resistant cells	AI923458		Williams Beuren syndrome chromosome region 21	Hs.182476	106	231	NP_112585
107	Resistant cells	NM_000597		insulin-like growth factor binding protein 2 (36kD)	Hs.43326	107	232	NP_000588
108	Resistant cells	U90304		iroquois homeobox protein 5	Hs.25351	108	233	P78411
109	Resistant cells	NM_004968		islet cell autoantigen 1 (69kD)	Hs.167927	109	234	NP_004959
110	Resistant cells	AL563283		androgen-induced basic leucine zipper	Hs.372924	110	235	NP_570968
111	Resistant cells	AA135522		KIAA0089 protein	Hs.82432	111	236	AAH28726
112	Resistant cells	AI867102	yes	solute carrier family 9 (sodium/hydrogen exchanger), isoform 3 regulatory factor 1	Hs.184276	112	237	XP_051621
113	Resistant cells	AW134976		KIAA0984 protein	Hs.11912	113	238	BAA76828
114	Resistant cells	AW665865		KIAA1069 protein	Hs.193143	114	239	BAA83021
115	Resistant cells	AB051487		nucleoporin 210	Hs.270404	115	240	BAB40814
116	Resistant cells	AB050049		methylcrotonoyl-Coenzyme A carboxylase 2 (beta)	Hs.167531	116	241	Q9HCC0
117	Resistant cells	NM_016835		microtubule-associated protein tau	Hs.101174	117	242	NP_058519
118	Resistant cells	AK002075		myelin gene expression factor 2	Hs.44268	118	243	NP_057216
119	Resistant cells	NM_000933		Homo sapiens mRNA; cDNA DKFZp434E235 (from clone DKFZp434E235)	Hs.348724	119	244	NP_000924
120	Resistant cells	AI435670		prostate epithelium-specific Ets transcription factor	Hs.79414	120	245	NP_036523
121	Resistant cells	NM_006443		putative c-Myc-responsive	Hs.109752	121	246	NP_006434
122	Resistant cells	AW263542		ESTs	Hs.403937	122		AAH15948

Gene No.	Highly Expressed in	Genbank Accession #	Modulated by BMS-A	Unigene Title	Unigene Cluster	DNA SEQ ID NO:	Amino Acid SEQ ID NO:	Protein ID
123	Resistant cells	AF153330		dual specificity phosphatase 16	Hs.20281	123	247	Q9BY84
124	Resistant cells	BC002702		solute carrier family 25 (mitochondrial carrier; ornithine transporter) member 15	Hs.78457	124	248	Q9Y619
125	Resistant cells	NM_006416		solute carrier family 35 (CMP-sialic acid transporter), member 1	Hs.82921	125	249	NP_006407
126	Resistant cells	NM_030674		solute carrier family 38, member 1	Hs.18272	126	250	NP_109599
127	Resistant cells	AF212371		spinster-like protein	Hs.379091	127	251	AAH08325
128	Resistant cells	AF096304		solute carrier family 19 (thiamine transporter), member 2	Hs.30246	128	252	AAD09765
129	Resistant cells	AK000948		trichorhinophalangeal syndrome 1	Hs.26102	129	253	Q9UHF7
130	Resistant cells	AI859834		ESTs, Moderately similar to hypothetical protein FLJ20489	Hs.445020	130		
131	Resistant cells	BF512846		ESTs	Hs.442762	131		
132	Resistant cells	NM_022969	yes	fibroblast growth factor receptor 2 (bacteria-expressed kinase, keratinocyte growth factor receptor, craniofacial dysostosis 1, Crouzon syndrome, Pfeiffer syndrome, Jackson-Weiss syndrome)	Hs.278581	132	254	NP_075258
133	Resistant cells	AA741493	yes	ESTs	Hs.143842	133		
134	Resistant cells	NM_001424		epithelial membrane protein 2	Hs.29191	134	255	P54851
135	Resistant cells	AW242920	yes	ESTs	Hs.129368	135		
136	Resistant cells	W44413		small protein effector 1 of Cdc42	Hs.22065	136	256	Q9HB17
137	Resistant cells	AK021717		Homo sapiens cDNA FLJ11655 fs, clone HEMBA1004554	Hs.287436	137		

Table 3 presents a resistance/sensitivity prediction of the 23 breast cell lines for BMS-A in the 'leave one out' cross validation test using a Weighted Voting algorithm. The true class is assigned as in Table 1, based on the IC₅₀ results. The predicted class was determined by using the optimal 15 and 7 polynucleotides as the predictor set to predict the resistance or sensitive class. These polynucleotides were selected either from the 137 polynucleotides derived from three analysis methods as shown in Table 2, or from 40 drug treatment modulated polynucleotides as indicated in Table 2. "S" represents Sensitive; "R" represents Resistant. The PS score refers to prediction strength for each prediction made on a cell line by the predictor set. The PS score ranges from 0 to 1, i.e., corresponding from low to high confidence in making the prediction. The error predictions are indicated by an asterisk (*).

TABLE 3

Cell Line	True Class	15 markers from 137 polynucleotides in Table 2			7 modulated markers from 40 polynucleotides as indicated in Table 2		
		Predicted Class	PS score	Error?	Predicted Class	PS score	Error?
MDAMB157	S	S	0.627		S	0.696	
MDAMB231	S	S	0.857		S	1.000	
HCC1954	S	S	0.416		S	0.847	
HCC70	S	S	0.695		S	1.000	
BT20	S	S	0.586		S	0.794	
HCC1806	S	S	0.985		S	1.000	
Hs578T	S	S	0.775		S	0.570	
HCC1419	R	R	1.000		R	1.000	
SkBr3	R	R	0.852		R	0.992	
AU565	R	R	0.629		R	0.763	
HCC38	R	S	0.101	*	S	0.501	*
BT474	R	R	0.938		R	1.000	
MDAMB468	R	R	0.392		R	0.416	
HCC1428	R	R	0.623		R	0.939	
MDAMB435S	R	S	0.723	*	R	0.324	
H3396	R	R	1.000		R	1.000	
BT549	R	R	0.029		R	0.012	
Zr-75-30	R	R	0.958		R	1.000	
MCF7	R	R	0.911		R	1.000	
Her2MCF7	R	R	0.991		R	1.000	
MDAMB436	R	R	0.340		R	0.412	
Zr-75-1	R	R	1.000		R	1.000	
MDAMB453	R	R	0.983		R	1.000	

Table 4 lists the predictor set of 15 polynucleotides used in prediction as shown in Table 3. These 15 polynucleotides were selected from the 137

polynucleotides derived from three analysis methods as shown in Table 2. The relative expression pattern, i.e., sensitive or resistant, polynucleotide description and Unigene cluster number for this 15 predictor polynucleotide subset are indicated in Table 4.

5

TABLE 4

Highly Expressed in:	Modulated by BMS-A	Unigene Title	Unigene Cluster No
Sensitive cells		BphB2	Hs.125124
Sensitive cells		parvin, alpha	Hs.44077
Sensitive cells	yes	coagulation factor II (thrombin) receptor-like 1	Hs.154299
Sensitive cells	yes	aldo-keto reductase family 1, member C3	Hs.78183
Sensitive cells	yes	interferon, gamma-inducible protein 16	Hs.155530
Sensitive cells	yes	jagged 1 (Alagille syndrome)	Hs.91143
Sensitive cells		hypothetical protein MGC13105	Hs.22744
Sensitive cells		snail homolog 2 (Drosophila)	Hs.93005
Resistant cells		Homo sapiens mRNA cDNA DKFZp761C1712	Hs.4774
Resistant cells	yes	Homo sapiens cDNA FLJ31753 fis, clone NT2RI2007468	Hs.349283
Resistant cells		ATP-binding cassette, sub-family A (ABC1), member 3	Hs.26630
Resistant cells		CGI-81 protein	Hs.279583
Resistant cells	yes	ESTs	Hs.445708
Resistant cells		EST	Hs.159643
Resistant cells		hypothetical protein LOC112868	Hs.97837

Table 5 lists the predictor set of 7 polynucleotides used in prediction as shown in Table 3. These 7 polynucleotides were selected from the 40 polynucleotides that were modulated by drug treatment as indicated in Table 2. The relative expression pattern, i.e., sensitive or resistant, polynucleotide description and Unigene cluster number for this 7 predictor polynucleotide subset are indicated in Table 5.

10

TABLE 5

Highly Expressed in:	Modulated by BMS-A	Unigene Title	Unigene Cluster No
Resistant cells	yes	Homo sapiens cDNA FLJ31753 fis, clone NT2RI2007468	Hs.349283
Sensitive cells	yes	jagged 1 (Alagille syndrome)	Hs.91143
Sensitive cells	yes	interferon, gamma-inducible protein 16	Hs.155530
Sensitive cells	yes	coagulation factor II (thrombin) receptor-like 1	Hs.154299
Resistant cells	yes	ESTs	Hs.445708
Sensitive cells	yes	aldo-keto reductase family 1, member C3	Hs.78183
Sensitive cells	yes	polymerase I and transcript release factor	Hs.29759

Table 6 lists the representative RT-PCR primer sets for each of the protein tyrosine kinase biomarker polynucleotides of the present invention. The SEQ ID NO: for each RT-PCR primer is provided (SEQ ID NO:257 thru 530).

5

TABLE 6

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
NM_004431	Forward Primer	TCCTCACACTAAGAGGGCAGA	257
NM_004431	Reverse Primer	ACCTCAACACAACCAAGCATC	258
AF025304	Forward Primer	TCAGTGAGTACAACGCCACAG	259
AF025304	Reverse Primer	CTTCTCCTGGATGCTTGTCTG	260
NM_001753	Forward Primer	CCACCTTCACTGTGACGAAAT	261
NM_001753	Reverse Primer	CCAGATGTGCAGGAAAGAGAG	262
NM_001233	Forward Primer	AGCTGTCTGCACATCTGGATT	263
NM_001233	Reverse Primer	CCTGGGGTCCAAGTATCAAT	264
NM_000700	Forward Primer	CATCAAGCCATGAAAGGTGTT	265
NM_000700	Reverse Primer	ACAAAGAGCCACCAGGATTTT	266
NM_004039	Forward Primer	GAAGTATGTTCCCAAGTGGGA	267
NM_004039	Reverse Primer	AACCAGGTTCAAGGAAAGCATT	268
BG107577	Forward Primer	TTTCGTGAACAAGCACCTGA	269
BG107577	Reverse Primer	ATGAGCTCAAAGGCAAAGGA	270
BE965369	Forward Primer	GTTTAAATCCGGATTGGCAT	271
BE965369	Reverse Primer	GTGGCCGTGATAATTTTGAA	272
NM_001993	Forward Primer	AAAATGGAAGGAAATTGGGTG	273
NM_001993	Reverse Primer	TGCCCAGAATACCAATGTCTC	274
BF792126	Forward Primer	TCGGTGAATTCAAGGACCAT	275
BF792126	Reverse Primer	GCTGCCTTCAAGGATCTCAC	276
E856341	Forward Primer	TGCCAGGTAAAGCTCTGTCC	277
E856341	Reverse Primer	GTCTGTGGATGAGCATGTG	278
U17496	Forward Primer	ATCTCCAGAGCTCGCTTTACC	279
U17496	Reverse Primer	TTCACCCGTAAGGCACTAATG	280
NM_002800	Forward Primer	TATGGTTATGTGGATGCAGCA	281
NM_002800	Reverse Primer	AGATGACTCGATGGTCCACAC	282
NM_000311	Forward Primer	CCGAGTAAGCCAAAAACCAA	283
NM_000311	Reverse Primer	CTCATCCATGGGCCTGTAGT	284
NM_003739	Forward Primer	GGTGAGGAACCTTCACCAACA	285
NM_003739	Reverse Primer	CTTGAGTCCTGGCTTGTGAG	286
NM_020639	Forward Primer	TACTTGGGTGAGTCCTTGTGG	287
NM_020639	Reverse Primer	GACTCTTAGGCCTGTGGCTCT	288
AF208043	Forward Primer	GGAGTAAGGTGTCCGAGGAAC	289
AF208043	Reverse Primer	CTGACATTTGGCCACTGTTT	290
AF003837	Forward Primer	CCTGTAACATAGCCCGAAACA	291
AF003837	Reverse Primer	AGTTGTCTCCATCCACACAGG	292
BC002832	Forward Primer	ACGTGTATGCAGATGGAAAGG	293
BC002832	Reverse Primer	CAGAGGCTGTGACGTTGTGTA	294
NM_006994	Forward Primer	AATTTGTGCAGTTGGGAGATG	295
NM_006994	Reverse Primer	TGATCTCTACCCTGCAGCTGT	296
AF327443	Forward Primer	CATCTGACTTCACCTGTGGGT	297
AF327443	Reverse Primer	TTCTGACTGTCCCTGCTGACT	298
NM_021615	Forward Primer	ACCCCGACGTCTTCTACCTAA	299

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
NM_021615	Reverse Primer	GCAGATAGGCATCAAACACGT	300
AF104857	Forward Primer	AGTTCCTGGGCATAATGAGT	301
AF104857	Reverse Primer	AACATGAGAGCTTGGGATCCT	302
AL136896	Forward Primer	AGCCGAATCCACTCTCATGT	303
AL136896	Reverse Primer	TAACAAGGCACAGCAAGCAG	304
AL565621	Forward Primer	CTCAGACCTTTGCCCTTCTCT	305
AL565621	Reverse Primer	TCCGGCTCAGACTGAATAAGA	306
BF111719	Forward Primer	CACACATGGGCATTGTCTTA	307
BF111719	Reverse Primer	GGATATGCAGTGGGAAGGAA	308
BC020608	Forward Primer	CACCGAGAATCCTTACACCAA	309
BC020608	Reverse Primer	CAGAATCCATCCTCCTTCCTC	310
AW575374	Forward Primer	CATGCACACACACAGAATG	311
AW575374	Reverse Primer	TTTCCTTTGGAACTGGGATT	312
NM_001401	Forward Primer	CTTGCTGAATTCAACTCTGCC	313
NM_001401	Reverse Primer	AAACCACAGAGTGGTCATTGC	314
BC001247	Forward Primer	AGGAGAAGGAAGACAAGCCAG	315
BC001247	Reverse Primer	CTTGCTGATTTCTGCTTCAGG	316
BE669858	Forward Primer	CTGCTTGAGACTGTTCTGGCT	317
BE669858	Reverse Primer	GATTAGAGGGCTTCCTCATGG	318
NM_000127	Forward Primer	CAAGGGGAAGAGGTACCTGAC	319
NM_000127	Reverse Primer	TCTGTACAGCGAGAATCCTT	320
NM_002589	Forward Primer	GACTCTGGGCGTCTCTGAAG	321
NM_002589	Reverse Primer	CAGCAACAAGCCAGTCTCAA	322
AI133452	Forward Primer	ACATCATGAGTTGGTCCTTGC	323
AI133452	Reverse Primer	AATCTGCAATGCCACAGGTAG	324
NM_006101	Forward Primer	TCCTCATACATGGCCTCACA	325
NM_006101	Reverse Primer	TGTCGGCACCCTCATAAAA	326
AL135264	Forward Primer	GGTGCAGGTTGACACTGAAA	327
AL135264	Reverse Primer	AAGGTTCAACAGGACACAGG	328
NM_014164	Forward Primer	ATCACAGGCATCATCCTC	329
NM_014164	Reverse Primer	GGTTGTGAGCTCCTGTTTCTG	330
BC003502	Forward Primer	GGGGTGTAGGTGGGAGTCAC	331
BC003502	Reverse Primer	AGTGCCCTCAGCCAAAATGT	332
AA780067	Forward Primer	GCCATCCTCTTGATAAGCTGA	333
AA780067	Reverse Primer	TCTTCCAGGATTCTCTTTGG	334
AA702248	Forward Primer	GATTGCAGATCCTATGCAGGA	335
AA702248	Reverse Primer	GCATCCAGGACAACACAAAGT	336
BC004372	Forward Primer	AAGGTGGAGCAAACACAACC	337
BC004372	Reverse Primer	TCCACTTGGCTTTCTGTCTT	338
BF688144	Forward Primer	CAAGTGCCCATTTAGGTTTGA	339
BF688144	Reverse Primer	ACTGACAGATGGCTCATTGG	340
NM_018067	Forward Primer	GAACACCAGAGACACTCCTGC	341
NM_018067	Reverse Primer	ACATCCTGGTAGGTGATGCAG	342
BG111761	Forward Primer	CGCATCTGTCCAGCATCTTA	343
BG111761	Reverse Primer	CAAAACCGGGACGCTAACT	344
NM_017821	Forward Primer	AGAAACAGTGGATCACGTTGG	345
NM_017821	Reverse Primer	TTCCAAGGGAATACCCAAAAC	346
AA722799	Forward Primer	GTTTCCACTTTTCCCAAGTGC	347
AA722799	Reverse Primer	TCACATGAAACGATTCTCTGCT	348
BC006436	Forward Primer	AATGTCAAAAAGTGTGGGCAAG	349
BC006436	Reverse Primer	ATGTGGACCGAGTAAAGGCTT	350
NM_006548	Forward Primer	CAGTCCCGGGTAGATATCCAT	351
NM_006548	Reverse Primer	TCTTCGGCTAGTTTGGTCTCA	352

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
NM_002194	Forward Primer	TGATTTGCCACAGTTGGTGTA	353
NM_002194	Reverse Primer	CTAGGTATGCGTCTCTGCAGG	354
BG251556	Forward Primer	CAGCCTGGTTTACAAATTCCA	355
BG251556	Reverse Primer	TGGGGAAAACTAAGGCAAAGT	356
J03202	Forward Primer	CAACAATGAAGCCTGCTCTTC	357
J03202	Reverse Primer	CCTGCTTCAGTGAGAGAATGG	358
NM_000245	Forward Primer	AGGACCGGTTTCATCAACTTCT	359
NM_000245	Reverse Primer	TCAATGTAGGACTGGTCCGTC	360
NM_002444	Forward Primer	AAATGGTGCCTTCAAGACCTT	361
NM_002444	Reverse Primer	CCGGCCTATACTCCTACAAGG	362
NM_012334	Forward Primer	TAATGGTGGTCTGAACAAGGC	363
NM_012334	Reverse Primer	AGTTGGCCCAAGTCCTTAAAA	364
AI769569	Forward Primer	CATGGAGGAGCCATACAACA	365
AI769569	Reverse Primer	TTTGTCTGCTCCCAAATTC	366
NM_002633	Forward Primer	TGCTTTGTATGAGACCCCAAC	367
NM_002633	Reverse Primer	CATCTTTCTCACGGATGTGGT	368
BC004295	Forward Primer	AGAAGACAGAGAGGTCAGCCC	369
BC004295	Reverse Primer	TGGGACCCTAATTTTCTGGAC	370
NM_016205	Forward Primer	ACCCTTGAGTTTTCGCCTCT	371
NM_016205	Reverse Primer	GGATCAAAGCAAAACCTGGA	372
NM_004815	Forward Primer	GCCCCTTTTGTATAGGACTGC	373
NM_004815	Reverse Primer	AATTCCAGTGAGGCACAAATG	374
NM_002872	Forward Primer	CAAGACCTGCCTTCTCATCAG	375
NM_002872	Reverse Primer	GAAGACGTCCGTCTGTGGATA	376
AF329267	Forward Primer	CAATTCTCTCAGCAGACCTGG	377
AF329267	Reverse Primer	ACCACGGAGTCAAAACCTTCT	378
AI572079	Forward Primer	CCCCAAGGCACATACTGTAA	379
AI572079	Reverse Primer	TGCCCATTTGTTGAACTAAAGC	380
NM_001549	Forward Primer	GAACATGCTGACCAAGCAGA	381
NM_001549	Reverse Primer	CAGTTGTGTCCACCCTTCCT	382
D50683	Forward Primer	AACAATACTGGCTGATCACCG	383
D50683	Reverse Primer	CATGGAGTGTGATCACTGTGG	384
NM_005902	Forward Primer	GGACTGCAGTGTGGAGTTCA	385
NM_005902	Reverse Primer	GAGAGGGGAGGGAGACAGAC	386
NM_014452	Forward Primer	GGTTTATAAGCCTTTGCCAGG	387
NM_014452	Reverse Primer	GTGGGAAAAGTCACACTGCAT	388
AB017644	Forward Primer	CTCCTCCTAATTGCAGTGCTG	389
AB017644	Reverse Primer	GTGATAGATTCTGGTGCGGAA	390
BC002323	Forward Primer	CCTCAGGTCCAACCTCATGT	391
BC002323	Reverse Primer	GTGCCCCAATTTTGTATTG	392
AL157452	Forward Primer	AGCCTTGTCTCCCTTGGATT	393
AL157452	Reverse Primer	TCAGTTGCCCTCTACAACC	394
BF752277	Forward Primer	AAGGCCCTGGATTCTCACTC	395
BF752277	Reverse Primer	GCCAGGACACCTTCAGAGAG	396
BF512299	Forward Primer	AAGAGCCTCCCAAAGGAAA	397
BF512299	Reverse Primer	GGGAAATGAAAGTGGCAAGA	398
AL049381	Forward Primer	TTGTTGGTTTTATTCTCCCC	399
AL049381	Reverse Primer	CAGTTGGAATCAAAGGGACA	400
NM_002585	Forward Primer	AGTGAGGAAGCCAAAGAGGAG	401
NM_002585	Reverse Primer	TTTGGCAGCATAAATATTGGC	402
T68445	Forward Primer	CAGAGAGGGAACCACCAGAG	403
T68445	Reverse Primer	CCCTGGGGAAATTTAAATGA	404
BF308645	Forward Primer	CTCTGTCGGGAAAGGAGAGA	405

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
BF308645	Reverse Primer	GAACTTTGACGACACCGACA	406
AF088867	Forward Primer	CTCTGGCCAGAGATACCACAG	407
AF088867	Reverse Primer	CATCAAGGGTTTGTGCTTGT	408
NM_004040	Forward Primer	AACTATGTGGCCGACATTGAG	409
NM_004040	Reverse Primer	CACCGAGAAGCACATGAGAAAT	410
AF151810	Forward Primer	CCTGAAGAACCGTGATGTCAT	411
AF151810	Reverse Primer	CTGTGCTCTGGATGAGGTAGC	412
NM_004252	Forward Primer	CACATCCCCTTCTTGACAAA	413
NM_004252	Reverse Primer	GATGAGGCACTCAGTGAGGAG	414
NM_005749	Forward Primer	TTGAAACCTAATTTTGTGGCG	415
NM_005749	Reverse Primer	AAATGTTGACACGTCTCCTGG	416
NM_003225	Forward Primer	CCTAATACCATCGACGTCCCT	417
NM_003225	Reverse Primer	AGCTCTGGGACTAATCACCGT	418
AA181060	Forward Primer	AAAAGGCTGACAACTGACCA	419
AA181060	Reverse Primer	TCACAGCCTAGGTAAGAGCCA	420
AL050025	Forward Primer	CAGGTACGAATTTTGCGGTTA	421
AL050025	Reverse Primer	TCGCAATCCACTCTCTGACTT	422
NM_001089	Forward Primer	CTCCTTCAGCTTCATGGTCAG	423
NM_001089	Reverse Primer	TCTGGCTCAGAGTCATCCAGT	424
NM_004915	Forward Primer	CAACCCAGCAGATTTTGTCTAT	425
NM_004915	Reverse Primer	CGAGGTCTCTCTTGTGGTCTG	426
AL523275	Forward Primer	TCTTTGCATTGAGATTGGTCC	427
AL523275	Reverse Primer	ACCGTGAAAAATGCACATCTC	428
NM_001218	Forward Primer	CCTTCAATCCGTCCTATGACA	429
NM_001218	Reverse Primer	GGAAGCAGCTCTTCAATGTTG	430
NM_016286	Forward Primer	GAGTGAATGCAGTAAACCCCA	431
NM_016286	Reverse Primer	CACTCAGCAGAAAGAGGATGG	432
BC000185	Forward Primer	CATCGAGGACGCTACTTCAAG	433
BC000185	Reverse Primer	AAAATAGGCCTGACGACACCT	434
NM_005505	Forward Primer	TTGGACAAACTGGGAAGATTG	435
NM_005505	Reverse Primer	ACGTAAGGGCATAGTGCATC	436
NM_016048	Forward Primer	GGGGATATTATTAGCGTGGGA	437
NM_016048	Reverse Primer	TGCCGCTTCTACTTCTGGTAA	438
BC000195	Forward Primer	TCCACTCACATTTCTATCGG	439
BC000195	Reverse Primer	GATTCCATTTACGGGGAAAAA	440
NM_001306	Forward Primer	AACCTGCATGGACTGTGAAAC	441
NM_001306	Reverse Primer	AATATCAAGTGCCCCTTCCAG	442
BC000021	Forward Primer	GCAAGTATAGCGCATTTGAGC	443
BC000021	Reverse Primer	CGTCTTGAAGTCCATGGAGAG	444
W68084	Forward Primer	TTAGATCTGAAGCCCTGGGTT	445
W68084	Reverse Primer	TGCTTGGTGAACATAACACCA	446
AA825563	Forward Primer	AGAAGAAAAACCCAAATGGCA	447
AA825563	Reverse Primer	TCCATAGTGGTTTTTACCAGCA	448
BE887449	Forward Primer	TGCGTACCAGGATTGGTTAAG	449
BE887449	Reverse Primer	GATGTCCAACAAAACGCTCAT	450
AI123815	Forward Primer	TGAGCATGGTATACTTTTGGG	451
AI123815	Reverse Primer	AAGCTTATAGGAATGGGCCAG	452
AI308862	Forward Primer	TGGGAAAAATTAAAACCCACA	453
AI308862	Reverse Primer	TCAAAGTGCCCTTTGGTAGTG	454
AW006352	Forward Primer	TCCTCAAACACAAAATCCCAG	455
AW006352	Reverse Primer	CTCCTACTATGGGCCTCCAAC	456
AL554277	Forward Primer	GAAGCAGATCGTCCTGAACTG	457
AL554277	Reverse Primer	GCTCATCATCCTCTTCTCCCT	458

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
BG289001	Forward Primer	TCCCAATAGCTTGTGGATCAG	459
BG289001	Reverse Primer	ATCAACCAGGAAGCCAACTTT	460
AI935915	Forward Primer	GACCAACACCTCTCCTAAGGG	461
AI935915	Reverse Primer	GTTGGGAGGGGACCATAGTTA	462
NM_017689	Forward Primer	GAAATAGCAAAAACAAGGCC	463
NM_017689	Reverse Primer	CAATGCAGCATGCTAGAAA	464
NM_017966	Forward Primer	CAGAATGTAAAGGGTGGGGAT	465
NM_017966	Reverse Primer	CCCTGAGACCTGGTTTACCTC	466
AI923458	Forward Primer	GATGGCAGCTATGAAGTCCTG	467
AI923458	Reverse Primer	GCATTCCAGCTATCACCTGAA	468
NM_000597	Forward Primer	CACCTCTACTCCCTGCACATC	469
NM_000597	Reverse Primer	AGAAGAGATGACACTCGGGGT	470
U90304	Forward Primer	TTTGGCTAAAGACCCGAAAAT	471
U90304	Reverse Primer	TCTCTCTCTCTCGGTGATGGA	472
NM_004968	Forward Primer	GAGCAGGAAAGATGATGCAAG	473
NM_004968	Reverse Primer	AAGTATCTGAGATGGCCCGAT	474
AL563283	Forward Primer	CTCTGGAATGGACTGAAGCTG	475
AL563283	Reverse Primer	AAAAGTCCAGGAGCTGGAGAG	476
AA135522	Forward Primer	CACCTCATCACAAACACCCTCT	477
AA135522	Reverse Primer	TGCTAGGATCCACCCTCCTAT	478
AI867102	Forward Primer	CTCTTCCCAGCTCCTGATTCT	479
AI867102	Reverse Primer	CTGAAGGACTGAAGGGAGCTT	480
AW134976	Forward Primer	ACATGCTGTGTGCTAGAGGCT	481
AW134976	Reverse Primer	AACATGCATGCATTGTACCAA	482
AW665865	Forward Primer	TTCCAGGAAGAACATCATTGC	483
AW665865	Reverse Primer	CTTTTCCTTCAGGGAACCAAG	484
AB051487	Forward Primer	TTCTCAGCCAAAGCAGATGTT	485
AB051487	Reverse Primer	TGCTTCTCCTCAGCAATTTGT	486
AB050049	Forward Primer	ACTATGGGATGTGTGGCAGAG	487
AB050049	Reverse Primer	GCTCTTTTAAAGCCGCTTCAT	488
NM_016835	Forward Primer	AAAGAGGCTGACCTTCCAGAG	489
NM_016835	Reverse Primer	AAGGCAAGGCCTATTTTCAA	490
AK002075	Forward Primer	GAAGCAATGAATAGCATGGGA	491
AK002075	Reverse Primer	CCATTCTCCAGTCACACTGT	492
NM_000933	Forward Primer	TCGGTCTTGGCTACTTGAAGA	493
NM_000933	Reverse Primer	CAGCGTTCCAGAAAATCTGAG	494
NM_012391	Forward Primer	AAGGAGTTGCTACTCAAGCCC	495
NM_012391	Reverse Primer	CTTGTAATACTGGCGGATGGA	496
NM_006443	Forward Primer	CCATCCTTGGGTGTAGGCTAT	497
NM_006443	Reverse Primer	CTCGAAGTATCGATCCAGCAG	498
BC015948	Forward Primer	ATGTGCCCTCACATCTGTTTC	499
BC015948	Reverse Primer	GGGTTTTAACAGCAGGGTAGC	500
AF153330	Forward Primer	GAAATCAGTCTACCAAGGGGC	501
AF153330	Reverse Primer	CGACTTTGCAATCTTGACACA	502
BC002702	Forward Primer	GAAGAGTGGGCAAACATGAAA	503
BC002702	Reverse Primer	CCCACCTGGGAGTAAGTCTTC	504
NM_006416	Forward Primer	CCAGGTGACCTACCAAGTTGAA	505
NM_006416	Reverse Primer	TTCCACCACCACTTTTGTAGC	506
NM_030674	Forward Primer	TGGCAAACACTGGAATCCTAC	507
NM_030674	Reverse Primer	TCTGTAGAGAGGTGGCTCCAA	508
AF212371	Forward Primer	CGGATGCTCAGCATCTTCTAC	509
AF212371	Reverse Primer	ACTACCAGGAACAGCAGCAGA	510
AF096304	Forward Primer	AGGCAATCCGATTACGACTT	511

Genbank Accession No.	RT-PCR Primer Type	Rt-PCR Primer Sequence	SEQ ID NO:
AF096304	Reverse Primer	CTCTGCCTCCTTCATCAACAG	512
AK000948	Forward Primer	AGAAGGACTTCTCCAGCAAGG	513
AK000948	Reverse Primer	CTGGGACAGAATGGACAGTGT	514
AI859834	Forward Primer	TGGCCATTTCAGACAGCATTA	515
AI859834	Reverse Primer	CAGTACTTGGGAGGCTGAG	516
BF512846	Forward Primer	GGGCCCCACTTGACTCATTTA	517
BF512846	Reverse Primer	GCCTGCAGAGATCTCACTTTG	518
NM_022969	Forward Primer	ACAGGATGGGCCTCTCTATGT	519
NM_022969	Reverse Primer	TCCTCAGGAACACGGTTAATG	520
AA741493	Forward Primer	ACACCTTGGTACCACCAATCA	521
AA741493	Reverse Primer	GGTCTCTTGCCTTCATCCAGT	522
NM_001424	Forward Primer	GCATCGCCTTCTTCATCTTC	523
NM_001424	Reverse Primer	CGTAGCTGCCTTCTCTGGTC	524
AW242920	Forward Primer	TTCATGCGTGAAAGTGTGAAG	525
AW242920	Reverse Primer	TTTGATCAAAGGGTGTATCAG	526
W44413	Forward Primer	GGTAGGGAGCTTCTCAGCAA	527
W44413	Reverse Primer	GTTAGCCCAGAGGAGCTCAA	528
AK021717	Forward Primer	CACAGAAAACACCCCCACTT	529
AK021717	Reverse Primer	ACTGTATGGAGGCCAGTTG	530

DETAILED DESCRIPTION OF THE INVENTION

The present invention describes the identification of polynucleotides that correlate with drug sensitivity or resistance of untreated cell lines to determine or predict sensitivity of the cells to a drug, compound, or biological agent. These polynucleotides, called marker or predictor polynucleotides herein, can be employed for predicting drug response. The marker polynucleotides have been determined in an *in vitro* assay employing microarray technology to monitor simultaneously the expression pattern of thousands of discrete polynucleotides in untreated cells, whose sensitivity to compounds or drugs, in particular, compounds that modulate, e.g., inhibit, protein tyrosine kinase or protein tyrosine kinase activity is tested. The protein tyrosine kinases, or activities thereof, associated with response to a drug, compound, or biological agent include, for example, members of the Src family of protein tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases. (See, e.g., P. Blume-Jensen and T. Hunter, 2001, "Oncopolynucleotide Kinase Signaling", *Nature*, 411:355-365).

The assay according to this invention has allowed the identification of the marker polynucleotides, called protein tyrosine kinase biomarkers herein, having

expression levels in the cells that are highly correlated with drug sensitivity exhibited by the cells. Such marker polynucleotides encompass the above-listed protein tyrosine kinase-encoding polynucleotides, and serve as useful molecular tools for predicting a response to drugs, compounds, biological agents, chemotherapeutic agents, and the like, preferably those drugs and compounds, and the like, that affect protein tyrosine kinase activity via direct or indirect inhibition or antagonism of the protein tyrosine kinase function or activity.

In its preferred aspect, the present invention describes polynucleotides that correlate with sensitivity or resistance of breast cell lines to treatment with a protein tyrosine kinase inhibitor compound, e.g., BMS-A, as described herein. (FIG. 1 and Table 2). The protein tyrosine kinase inhibitor compound, BMS-A, utilized for identifying the polynucleotide predictor sets of this invention, was described in WO 00/62778, published October 26, 2000, and is hereby incorporated by reference in its entirety. BMS-A has potent inhibitory activity for a number of protein tyrosine kinases, for example, members of the Src family of protein tyrosine kinases, including Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as the Bcr-abl, Jak, PDGFR, c-kit and Eph receptors protein tyrosine kinases. Specifically, for the BMS-A protein tyrosine kinase inhibitor compound analyzed, the expression of 137 predictor polynucleotides was found to correlate with resistance/sensitivity of the breast cell lines to the compound.

In accordance with the invention, an approach has been discovered in which polynucleotides and combinations of polynucleotides have been identified whose expression pattern, in a subset of cell lines, correlates to and can be used as an *in vitro* predictor of cellular response to treatment or therapy with one compound, or with a combination or series of compounds, that are known to inhibit or activate the function of a protein, enzyme, or molecule (e.g., a receptor) that is directly or indirectly involved in cell proliferation, cell responses to external stimuli, (such as ligand binding), or signal transduction, e.g., a protein tyrosine kinase. Preferred are antagonists or inhibitors of the function of a given protein, e.g., a tyrosine kinase.

In a preferred aspect, the BMS-A protein tyrosine kinase inhibitor was employed to determine drug sensitivity in a panel of breast cell lines following exposure of the cells to this compound. Some of the cell lines were determined to be

resistant to treatment with the inhibitor compound, while others were determined to be sensitive to the inhibitor. (Table 1). A subset of the cell lines examined provided an expression pattern or profile of polynucleotides, and combinations of polynucleotides, that correlated to, and thus serve as a predictor of, a response by the
5 cells to the inhibitor compound, and to compounds having similar modes of action and/or structure. (Figure 1 and Tables 2 and 4-5).

Such a predictor set of cellular polynucleotide expression patterns correlating with sensitivity or resistance of cells following exposure of the cells to a drug, or a combination of drugs, provides a useful tool for screening a cancer, tumor, or patient
10 test sample before treatment with the drug or a drug combination. The screening technique allows a prediction of cells of a cancer, tumor, or test sample exposed to a drug, or a combination of drugs, based on the polynucleotide expression results of the predictor set, as to whether or not the cancer, tumor, or test sample, and hence a patient harboring the cancer and/or tumor, will or will not respond to treatment with
15 the drug or drug combination. In addition, the predictor polynucleotides or predictor polynucleotide set can also be utilized as described herein for monitoring the progress of disease treatment or therapy in those patients undergoing treatment involving a protein tyrosine kinase, e.g., src tyrosine kinase, inhibitor compound or chemotherapeutic agent for a disease, e.g., breast cancer.

20 According to a particular embodiment of the present invention, oligonucleotide microarrays were utilized to measure the expression levels of over 44,792 probe sets in a panel of 23 untreated breast cell lines for which the drug sensitivity to the protein tyrosine kinase inhibitor compound was determined. This analysis was performed to determine whether the polynucleotide expression
25 signatures of untreated cells were sufficient for the prediction of chemosensitivity. Data analysis allowed the identification of marker polynucleotides whose expression levels were found to be highly correlated with drug sensitivity. In addition, the treatment of cells with the BMS-A protein tyrosine kinase inhibitor compound also provided polynucleotide expression signatures predictive of sensitivity to the
30 compound. Thus, in one of its embodiments, the present invention provides these polynucleotides, i.e., polynucleotide "markers" or "biomarkers" or "predictors", which show utility in predicting drug response upon treatment or exposure of cells to a drug.

In particular, the marker or predictor polynucleotides are protein tyrosine kinase biomarkers/polynucleotides encoding protein tyrosine kinase biomarker proteins/polypeptides, such as a src tyrosine kinase inhibitor biomarker.

5 The performance of the polynucleotide expression and marker polynucleotide identification analyses embraced by the present invention is described in further detail and without limitation herein below.

IC₅₀ Determination and Phenotype Classification Based on Sensitivity of Twenty-three Breast Cell Lines to Src tyrosine kinase Inhibitor Compounds

20 Twenty-three breast cell lines were treated with a protein tyrosine kinase inhibitor compound (i.e., BMS-A) to determine the individual IC₅₀ value. The average IC₅₀ values, along with standard deviations, were calculated from 2 to 5 individual determinations for each cell line. As shown in Table 1, a large variation in the IC₅₀ values (>1000-fold) was observed for the compound among the twenty-three breast cell lines.

15 The IC₅₀ value for each cell line was log₁₀ transformed. The mean of log₁₀(IC₅₀) across the twenty-three breast cell lines was calculated for the compound. The log₁₀(IC₅₀) for each cell line was normalized to the mean of log₁₀(IC₅₀) across the twenty-three breast cell lines for the compound. The cell lines with a log₁₀(IC₅₀) below the mean of log₁₀(IC₅₀) were classified as sensitive to the compound, and those
20 with a log₁₀(IC₅₀) above the mean of log₁₀(IC₅₀) were classified as resistant. Table 1 presents the resistance/sensitivity classifications of the twenty-three breast cell lines to the BMS-A compound. As observed in Table 1, seven cell lines were classified as sensitive and sixteen cell lines were classified as resistant to the protein tyrosine kinase inhibitor compound.

25 Identifying Polynucleotides that Significantly Correlated with Drug Resistance/Sensitivity Classification

Expression profiling data of 44,792 probe sets represented on the HG-U133 array set for twenty-three untreated breast cell lines were obtained and preprocessed as described in Example 1, Methods. The preprocessed data containing 5322
30 polynucleotides were analyzed using K-mean Nearest Neighborhood (KNN) algorithm and "signal to noise model" (T.R. Golub et al., 1999, Science, 286:531-537) to identify polynucleotides whose expression patterns were strongly correlated with

the drug resistance/sensitivity classification (Table 1). An "idealized expression pattern" corresponds to a polynucleotide that is uniformly high in one class (e.g., sensitive) and uniformly low in the other class (e.g., resistant). Initially, a KNN analysis was performed in which a correlation coefficient was obtained for each polynucleotide using "signal to noise model". The correlation coefficient, which is a measure of relative classification separation, is obtained using the following formula:

$$P(g,c)=(\mu_1 - \mu_2) / (\sigma_1 + \sigma_2).$$

In the above formula, for $P(g,c)$, P represents correlation coefficient between expression for gene, g , and the sensitivity/resistance classification, c ; μ_1 represents the mean polynucleotide expression level of samples in class 1; μ_2 represents the mean polynucleotide expression level of samples in class 2; σ_1 represents the standard deviation of polynucleotide expression for samples in class 1; and σ_2 represents the standard deviation of polynucleotide expression for samples in class 2.

Large values of $P(g,c)$ indicate a strong correlation between polynucleotide expression and resistance/sensitivity classification. When the correlation is compared to that of a random permutation test (randomly assigned classification), a significance measurement p -value is obtained. Then, the polynucleotides can be ranked according to the correlation coefficient obtained from this analysis, with the highest value indicating the best correlation of polynucleotide expression level with the resistance/sensitivity classification to the protein tyrosine kinase inhibitor compound in the twenty-three breast cell lines.

The KNN analysis demonstrated that hundreds of polynucleotides correlated to the drug resistance/sensitivity classification for the compound. Therefore, for greater stringency, three different methods were applied to select a smaller subset of polynucleotides that correlated with the drug resistance/sensitive classification for the compound:

First, a permutation test was performed to calculate the significance of the correlation coefficients obtained in the above-described KNN analysis. 350 polynucleotides whose ' p ' value was less than or equal to 0.01 were selected. Second,

the Pearson correlation coefficient (a dimensionless index that ranges from -1.0 to 1.0), was calculated, in which the IC₅₀ data were considered as a continuous variable and a linear regression model was utilized to correlate polynucleotide expression level with IC₅₀ values for the twenty-three breast cell lines. Those polynucleotides with a correlation coefficient greater than 0.35 or less than -0.35 were selected ($p < 0.05$). Finally, Welch t-test was performed, the polynucleotides with p-values equal to or less than 0.05 were selected.

When the three analyses were performed to select polynucleotides correlated with the drug resistance/sensitivity classification for compound BMS-A, the polynucleotide lists from the three analysis methods were obtained and compared. It was observed that there were 168 polynucleotides overlapped from the three analyses. Of these, 32 polynucleotides were redundantly represented more than once on the 168 polynucleotide list, and removed to just leaving one copy per unique gene. Therefore, 137 unique polynucleotides are identified and listed in Table 2. There are 68 polynucleotides highly expressed in the cell lines that were classified as sensitive to BMS-A, while 69 polynucleotides are highly expressed in the cell lines that were classified as resistant to BMS-A. Examples of the polynucleotides include caveolin-1, caveolin-2, and annexin A1 and annexin A2, which are substrates for src tyrosine kinase (M.T. Brown and J.A. Cooper, 1996, *Biochemica et Biophysica Acta*, 1287:121-149). EphA2 and EphB2 are tyrosine kinase receptors, they have diverse roles in carcinopolynucleotidesis (M. Nakamoto and A. D. Bergemann, 2002, *Microscopy Research and Technique* 59:58-67).

Identification of polynucleotides modulated by drug treatment

To identify polynucleotides regulated by a protein tyrosine kinase inhibitor compound, e.g., BMS-A, 11 breast cell lines (indicated in bold in the Table 1) having an IC₅₀ ranging from 0.0055 to 9.5 μ M were used in a drug treatment study. Cells were treated with or without the BMS-A compound (0.4 μ M) in 0.1% DMSO for 24 hours. Expression profiling was performed, and the data were analyzed using GeneChip[®] Expression Analysis software MAS 5.0 (Affymetrix, Santa Clara, CA). The polynucleotide expression of a cell line treated with drug was compared pairwise to the polynucleotide expression of the same cell line without drug treatment.

A change in p-value was calculated, indicating an increase, decrease or no change in polynucleotide expression. When the p-value was less than 0.0025, the change was considered to be significant. Analysis was performed for all 11 cell lines to compare the polynucleotide expression with and without drug treatment.

5 In addition, a pair-wise t-test with permutation analysis was applied. Polynucleotides that were significantly modulated by the drug treatment in sensitive cell lines and/or in resistant cell lines were identified. Polynucleotides whose expression was significantly changed in at least 3 cell lines were considered to be modulated by the drug. The polynucleotides, whose expression was significantly
10 correlated with drug resistance/sensitivity classification and modulated by drug treatment as well, are indicated in Table 2. Examples of such polynucleotides include EphA2 and caveolin-2, which were highly expressed in sensitive cells and were down regulated by treatment with the protein tyrosine kinase inhibitor compound BMS-A only in sensitive cell lines as shown in Figure 2.

15 Down regulation of the marker polynucleotides by the protein tyrosine kinase inhibitor compound treatment is also seen in PC3 prostate cell line which is tested to be very sensitive to BMS-A. As illustrated in Figure 3, a dose and time dependent polynucleotide expression decrease of EphA2 and caveolin-2 is observed when compared to the untreated control.

20 Since EphA2 belongs to family of tyrosine kinase receptors, it is possible to test whether cells treated with the protein tyrosine kinase inhibitor compound BMS-A would affect phosphorylation status of EphA2. Immunoblot analysis of protein level and phosphorylation status of EphA2 in nine breast tumor cell lines is shown in Figure 4. Cells were treated with 0.1 μ M of BMS-A for one hour. Cell lysates were
25 immuno-precipitated with EphA2 antibody and blot with EphA2 antibody or anti-phosphotyrosine antibody. The results indicate that EphA2 protein level does not change upon the drug treatment for one hour, but the phosphorylation at tyrosine residue is dramatically decreased with the drug treatment. Recombinant human EphA2 protein was also tested in an *in vitro* kinase assay and showed auto
30 dephosphorylation upon the protein tyrosine kinase inhibitor compound BMS-A treatment with an inhibitory IC₅₀ of 17 nM.

The identification of those polynucleotides whose expression levels are not only correlated with the sensitivity or resistance of breast cell lines to treatment with a protein tyrosine kinase inhibitor compound (e.g., BMS-A), but also differentially regulated or modified by treatment with the compound can provide additional information about biological function or activity. The expression levels of these polynucleotides are regulated, or their phosphorylation level is modulated by the inhibitor compound indicating these polynucleotides are likely to be directly or indirectly involved in one or more protein tyrosine kinase signaling pathways, for example, protein tyrosine kinases that are members of the Src family of tyrosine kinases, including Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors.

Utility of highly correlated polynucleotides to make predictions

Polynucleotides that correlate to a specific property of a biological system can be used to make predictions about that biological system and other biological systems. The Genecluster software or other programs can be used to select polynucleotides and combinations of polynucleotides that can predict properties using a "weighted-voting cross-validation algorithm" (T.R. Golub et al., 1999, *Science*, 286:531-537). In particular, the Genecluster software was used to build predictors that demonstrate the utility of polynucleotides that correlate to drug sensitivity and resistance. As used herein, the terms "predictor" or "predictor sets" are used as follows: a predictor or a predictor set refers to a single gene, or combination of polynucleotides, whose expression pattern or properties can be used to make predictions, with different error rates, about a property or characteristic of any given biological system.

The ability of polynucleotide expression patterns to predict a resistance/sensitive classification was further investigated using a "weighted-voting cross-validation algorithm" which uses a leave one out cross-validation strategy as described by T.R. Golub et al., 1999, *Science*, 286:531-537. The program was formatted to select the optimal number of polynucleotides whose expression pattern could be used to predict, with optimal accuracy, the classification of a cell line based on resistance or sensitivity toward a given protein tyrosine kinase inhibitor compound, e.g., BMS-A. A brief description of the cross-validation strategy of the program is described.

Based on the leave-one-out cross-validation strategy, a total of twenty-three prediction analyses (i.e., the number of cell lines in the data set) were performed in an iterative manner and the results of all twenty-three prediction analyses were combined to select the optimal number of polynucleotides that had optimal predictive accuracy.

5 In each separate prediction analysis, one cell line was withheld from the data set, and an optimal number polynucleotide predictor was built, based on the remaining twenty-two cell lines, and was subsequently used to predict the class of the withheld sample.

Figure 5 shows the real error rates using different numbers of polynucleotides in the predictor set and using different selections and combinations of markers for
10 predicting classes among the breast cell lines which were either resistant or sensitive to BMS-A. When the predictor sets were selected from the 137 polynucleotides as shown in Table 2, the lowest error rate of 6.3% was achieved in the cross-validation tests with 15 markers. Another predictor set comprised of 7 different polynucleotides selected from the 40 polynucleotides that were modulated by the drug treatment
15 achieved an error rate of 3.1%. This result indicates that polynucleotides which are not only correlated with drug sensitivity, but also modulated by drug, can provide a better and more accurate prediction in a predictor set.

The real error rates for predicting the sensitivity class of breast cell lines to BMS-A were compared with the real error rates using the same number of
20 polynucleotides as the predictor set in 20 cases in which classification for the breast cell lines was randomly assigned. As shown in FIG. 6, in the cross-validation tests, when the predictor set contained either 7 or 15 polynucleotides selected from different polynucleotide groups, the error rate for predicting sensitivity of BMS-A in the 23 breast cell lines was 3.1% and 6.3%, respectively. By contrast, the real error rates
25 ranged from 30% to 83% when using same number of polynucleotides for the predictor set in 20 cases in which classification for the breast cell lines was randomly assigned. This result demonstrated that the error rate value for predicting sensitivity to BMS-A in 23 breast cell lines was significantly lower than the error rate for predicting randomly assigned classification.

30 Table 3 shows the prediction accuracy of the optimal 15 and 7 polynucleotide predictor sets for the resistance/sensitive classification of the twenty-three breast cell lines to BMS-A in the leave-one-out cross validation tests. When a 15 polynucleotide

predictor set selected from the 137 polynucleotides which were derived from above mentioned three analysis methods (i.e., KNN, Pearson correlation between polynucleotide expression level and IC₅₀ values for the twenty-three breast cell lines, and t-test) was used in a leave-one-out cross-validation test, twenty-one out of twenty-
5 three samples were correctly predicted and two resistant cell lines, HCC38 and MDA-MB-435S were predicted to be sensitive to BMS-A. This resulted in a 6.3% real error rate, calculated as follows:

$$10 \quad \frac{(2/16 \text{ resistant} + 0/7 \text{ sensitive}) \times 100\%}{2}$$

When a 7 polynucleotide predictor set, selected from the 40 drug treatment modulated polynucleotides that were part of the 137 polynucleotides in Table 2, was used in a leave-one-out cross-validation test, only one resistant cell line, HCC38, was predicted to be sensitive to BMS-A. This resulted in a 3.1% real error rate, calculated
15 as follows:

$$\frac{(1/16 \text{ resistant} + 0/7 \text{ sensitive}) \times 100\%}{2}$$

20 In addition, a Prediction Strength ("PS") score for each prediction made on a cell line by the predictor set can be obtained from the Genecluster software. The "PS" score ranges from 0 to 1, measuring the margin of victory in each prediction using weighted-voting cross-validation algorithm (see, e.g., T.R. Golub et al., 1999, *Science*, 286:531-537). The higher the value of a PS score is, the more confident the
25 prediction make. The PS score values for each cell line using the optimal 15 or 7 polynucleotide predictor set, obtained as described above for BMS-A, are shown in Table 3. Note that even though the cell line BT549 was predicted correct to be resistant with both the 15 and 7 polynucleotide predictor sets, the PS score was very low, which means the confidence of prediction is low.

30 It will be appreciated that the exact number of polynucleotides that should comprise an optimal predictor set is not particularly established or defined. It is unlikely in the real world that any predictor set can be obtained with 100% or absolute

accuracy. This is due to the fact that there is a trade-off between the amount of additional information and robustness that are gained by adding more polynucleotides, and the amount of noise that is concomitantly added. In accordance with the present invention, different numbers of polynucleotides were tested in the predictor sets; data
5 were obtained and analyzed for a protein tyrosine kinase inhibitor, BMS-A. The selection of marker polynucleotides for use in the prediction set was well within the total number of polynucleotides, as shown in Table 2, that strongly correlated with the sensitivity class distinction.

Thus, in accordance with the present invention, an approach has been
10 developed in which polynucleotides and combinations of polynucleotides have been discovered whose expression pattern in a subset of cell lines correlates with, and can be used as a predictor of, response to treatment with compounds that inhibit the function of protein tyrosine kinases.

Predictor sets, error rates and algorithms used to demonstrate utility

15 The number of polynucleotides in any given predictor or predictor set may influence the error rate of the predictor set in cross validation experiments and with other mathematical algorithms. The data show that the error rate of a predictor is somewhat dependent on the number of polynucleotides in the predictor set and the contribution of each individual polynucleotide in the given predictor set and the
20 number of cell lines that are tested in the cross validation experiment. For example, in a given predictor set, one polynucleotide may contribute more significantly than other polynucleotides to the prediction.

It is very likely that if a polynucleotide significantly contributes to a predictor set, then it can be used in different combinations with other polynucleotides to
25 achieve different error rates in different predictor sets. For example, polynucleotide A alone gives an error rate of 30%. In combination with polynucleotides, B, C and D, the error rate becomes 10%; in combination with polynucleotides B, D and E, the error rate becomes 12%; while a combination of polynucleotide A with polynucleotides E-X gives an error rate of 8%, and so on. As demonstrated in FIG. 5,
30 different selection and combination of polynucleotides in a predictor set achieve different error rates in the cross-validation tests.

When the predictor sets were selected from the 137 polynucleotides as shown in Table 2, the lowest error rate of 6.3% was achieved in the cross-validation test with 15 markers as shown in Table 4. Another predictor set comprised of 7 polynucleotides (Table 5) selected from the 40 polynucleotides that were modulated by the drug treatment, achieved an error rate of 3.1%. This result indicates that polynucleotides which are not only correlated with drug sensitivity, but also modulated by the drug, can provide a better and more accurate prediction in a predictor set.

The error rates as described herein apply to the set of cell lines used in the cross-validation experiment. If a different set is used, or more cell lines are added to the original set tested, then different error rates may be obtained as described and understood by the skilled practitioner. Importantly, different combinations of polynucleotides that correlate to drug sensitivity can be used to build predictors with different prediction accuracy.

Expression pattern of the protein tyrosine kinase biomarkers in primary breast tumors

One hundred thirty-four primary breast tumor biopsies were obtained from clinic, and expression profiles of these samples were performed. The expression pattern of the 137 polynucleotides, that are highly correlated with a resistance/sensitivity phenotype classification of the 23 breast cell lines for the protein tyrosine kinase inhibitor compound BMS-A according to the present invention (as shown in FIG.1 and Table 2), were examined in the 134 primary breast tumors as demonstrated in Figure 7. Each row corresponds to a gene, with the columns corresponding to expression level in the different breast tumor samples. The individual polynucleotide encoding the protein tyrosine kinase biomarkers of the invention is in the same order as indicated in Figure 1. It is clear as shown in Figure 7 that a group of primary breast tumors (as indicated by the arrow) highly express the sensitive biomarkers of protein tyrosine kinase inhibitor of the invention. By contrast, another different group primary breast tumors highly express the resistant biomarkers. Although, whether these group of primary breast tumors highly expressing the sensitive biomarkers are really sensitive to the protein tyrosine kinase compounds, e.g., BMS-A is unknown and need to be tested, the fact that the primary breast tumors

exist similar expression pattern of the protein tyrosine kinase biomarkers as the sensitive breast cell lines gives a promise clue.

Applications of predictor sets

Predictor sets with different error rates can be used in different applications.

5 Predictor sets can be built from any combination of the polynucleotides listed in Table 2, or the predictor polynucleotide subsets of 15 and 7 polynucleotides, as presented in each of Tables 4 and 5, respectively, to make predictions about the likely effect of protein tyrosine modulator compounds, e.g., inhibitors, or compounds that affect a protein tyrosine kinase signaling pathway in different biological systems. The various
10 predictor sets described herein, or the combination of these predictor sets with other polynucleotides or other co-variants of these polynucleotides, can have broad utility. For example, the predictor sets can be used as diagnostic or prognostic indicators in disease management; they can be used to predict how patients with cancer might respond to therapeutic intervention with compounds that modulate the protein tyrosine
15 kinase family (e.g., the src tyrosine kinase family); and they can be used to predict how patients might respond to therapeutic intervention that modulate signaling through an entire protein tyrosine kinase regulatory pathway, such as, for example, the src tyrosine kinase regulatory pathway.

While the data described herein were generated in cell lines that are routinely
20 used to screen for and identify compounds that have potential utility for cancer therapy, the predictors can have both diagnostic and prognostic value in other diseases areas in which signaling through a protein tyrosine kinase or a protein tyrosine kinase pathway is of importance, e.g., in immunology, or in cancers or tumors in which cell signaling and/or proliferation controls have gone awry. Such protein tyrosine kinases
25 and their pathways comprise, for example, members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Although the data described herein have been generated using the particularly exemplified protein tyrosine kinase inhibitor compound, BMS-A, three other protein
30 tyrosine kinase inhibitor compounds were tested in addition to BMS-A and were found to have similar sensitivity and resistance classifications in the 23 breast cell lines evaluated. Thus, the predictors can have both diagnostic and prognostic value

related to other inhibitor molecules, as well as any molecules or therapeutic interventions that affect protein tyrosine kinases, such as Src tyrosine kinase, or a protein tyrosine kinase signaling pathways, such as that of the Src tyrosine kinase.

Those having skill in the pertinent art will appreciate that protein tyrosine
5 kinase pathways, e.g., the Src tyrosine kinase pathway, are present and functional in cell types other than cell lines of breast tissue. Therefore, the described predictor set of polynucleotides, or combinations of polynucleotides within the predictor set, can show utility for predicting drug sensitivity or resistance to compounds that interact with, or inhibit, a protein tyrosine kinase activity in cells from other tissues or organs
10 associated with a disease state, or cancers or tumors derived from other tissue or organ types. Non-limiting examples of such cells, tissues and organs include colon, breast, lung, heart, prostate, testes, ovaries, cervix, esophagus, pancreas, spleen, liver, kidney, intestine, stomach, lymphocytic and brain, thereby providing a broad and advantageous applicability to the predictor polynucleotide sets described herein. Cells
15 for analysis can be obtained by conventional procedures as known in the art, for example, tissue or organ biopsy, aspiration, sloughed cells, e.g., colonocytes, clinical or medical tissue, or cell sampling procedures.

Functionality of polynucleotides that make up a predictor set

The use of a predictor, or predictor set, (e.g., predictor polynucleotides, or a
20 predictor set of polynucleotides) allows for the prediction of an outcome prior to having any knowledge about a biological system. Essentially, a predictor can be considered to be a tool that is useful in predicting the phenotype that is used to classify the biological system. In the specific embodiment provided by the present invention, the classification as "resistant" or "sensitive" is based on the IC_{50} value of
25 each cell line to a compound (e.g., the protein tyrosine kinase inhibitor compound BMS-A as exemplified herein), relative to the mean $\log_{10}(IC_{50})$ value of the cell line panel (e.g., a twenty-three breast cell line panel, as exemplified herein).

As a particular example, a number of the polynucleotides described herein (Table 2) are known to be substrates for the src tyrosine kinase family, e.g., caveolin-1
30 and caveolin-2 (M.T. Brown and J.A. Cooper, 1996, *Biochemica et Biophysica Acta*, 1287:121-149). EphA2 is a tyrosine kinase receptor. The data presented herein demonstrated that EphA2 is highly expressed in the sensitive cell lines, and its

expression level and activity are down regulated by treatment of the protein tyrosine kinase inhibitor compound BMS-A. This is expected, since polynucleotides that contribute to a high predictor accuracy are likely to play a functional role in the pathway that is being modulated. For example, Herceptin therapy (i.e., antibody that
5 binds to the Her2 receptor and prevents function via internalization) is indicated when the Her2 polynucleotide is overexpressed. It is unlikely, although not impossible, that a therapy will have a therapeutic effect if the target enzyme is not expressed.

However, although the complete function of all of the polynucleotides and their functional products (proteins and mRNAs) that make up a predictor set are not
10 currently known, some of the polynucleotides are likely to be directly or indirectly involved in a protein tyrosine kinase signaling pathway, such as the Src tyrosine kinase signaling pathway. In addition, some of the polynucleotides in the predictor set may function in the metabolic or other resistance pathways specific to the compounds being tested. Notwithstanding, a knowledge about the function of the polynucleotides
15 is not a requisite for determining the accuracy of a predictor according to the practice of the present invention.

As described herein, polynucleotides have been discovered that correlate to the relative intrinsic sensitivity or resistance of breast cell lines to treatment with compounds that interact with and inhibit protein tyrosine kinases, e.g., Src tyrosine
20 kinase. These polynucleotides have been shown, through a weighted voting, leave-one-out, cross validation program, to have utility in predicting the intrinsic resistance and sensitivity of breast cell lines to these compounds.

An embodiment of the present invention relates to a method of determining or predicting if an individual requiring drug or chemotherapeutic treatment or therapy for
25 a disease, for example, a breast cancer or a breast tumor, will be likely to successfully respond or not respond to the drug or chemotherapeutic agent prior to subjecting the individual to such treatment or chemotherapy. The drug or chemotherapeutic agent can be one that modulates a protein tyrosine kinase activity or signaling involving a protein tyrosine kinase. Nonlimiting examples of such protein tyrosine kinases
30 include members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. In accordance with the method of the

invention, cells from a tissue or organ associated with disease, e.g., a patient biopsy of a tumor or cancer, preferably a breast cancer or tumor biopsy, are subjected to an *in vitro* assay as described herein, to determine their marker polynucleotide expression pattern (polynucleotides from Table 2 and/or the predictor polynucleotide subsets of
5 Tables 4-5) prior to their treatment with the compound or drug, preferably an inhibitor of a protein tyrosine kinase. The resulting polynucleotide expression profile of the cells before drug treatment is compared with the polynucleotide expression pattern of the same polynucleotides in cells that are either resistant or sensitive to the drug or compound, as provided by the present invention.

10 In another related embodiment, the present invention includes a method of predicting, prognosing, diagnosing, and/or determining whether an individual requiring drug therapy for a disease state or chemotherapeutic for cancer (e.g., breast cancer) will or will not respond to treatment prior to administration of treatment. The treatment or therapy preferably involves a protein tyrosine kinase modulating agent,
15 compound, or drug, for example, an inhibitor of the protein tyrosine kinase activity. Protein tyrosine kinases include, without limitation, members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Preferred is src tyrosine kinase and inhibitors thereof. In accordance with
20 this embodiment, cells from a patient's tissue sample, e.g., a breast tumor or cancer biopsy, are assayed to determine their polynucleotide expression pattern prior to treatment with the protein tyrosine kinase modulating agent, compound, or drug. The resulting polynucleotide expression profile of the test cells before exposure to the compound or drug is compared with that of one or more of the predictor subsets of
25 polynucleotides comprising either 15 or 7 polynucleotides as described herein and shown in Tables 4-5, respectively.

Success or failure of treatment of a patient's cancer or tumor with the drug can be determined based on the polynucleotide expression pattern of the patient's cells being tested, compared with the polynucleotide expression pattern of the predictor
30 polynucleotides in the resistant or sensitive panel of that have been exposed to the drug or compound and subjected to the predictor polynucleotide analysis detailed herein. Thus, if following exposure to the drug, the test cells show a polynucleotide

expression pattern corresponding to that of the predictor polynucleotide set of the control panel of cells that is sensitive to the drug or compound, it is highly likely or predicted that the individual's cancer or tumor will respond favorably to treatment with the drug or compound. By contrast, if, after drug exposure, the test cells show a polynucleotide expression pattern corresponding to that of the predictor polynucleotide set of the control panel of cells that is resistant to the drug or compound, it is highly likely or predicted that the individual's cancer or tumor will not respond to treatment with the drug or compound.

In a related embodiment, screening assays are provided for determining if a patient's cancer or tumor is or will be susceptible or resistant to treatment with a drug or compound, particularly, a drug or compound directly or indirectly involved in protein tyrosine kinase activity or a protein tyrosine kinase pathway, e.g., the Src tyrosine kinase activity or pathway.

Also provided by the present invention are monitoring assays to monitor the progress of a drug treatment involving drugs or compounds that interact with or inhibit protein tyrosine kinase activity. Protein tyrosine kinases encompassed by these monitoring assays include members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Such *in vitro* assays are capable of monitoring the treatment of a patient having a disease treatable by a compound or agent that modulates or interacts with a protein tyrosine kinase by comparing the resistance or sensitivity polynucleotide expression pattern of cells from a patient tissue sample, e.g., a tumor or cancer biopsy, preferably a breast cancer or tumor sample, prior to treatment with a drug or compound that inhibits the protein tyrosine kinase activity and again following treatment with the drug or compound with the expression pattern of one or more of the predictor polynucleotide sets described, or combinations thereof. Isolated cells from the patient are assayed to determine their polynucleotide expression pattern before and after exposure to a compound or drug, preferably a protein tyrosine kinase inhibitor, to determine if a change of the polynucleotide expression profile has occurred so as to warrant treatment with another drug or agent, or discontinuing current treatment. The resulting polynucleotide expression profile of the cells tested before and after

treatment is compared with the polynucleotide expression pattern of the predictor set of polynucleotides that have been described and shown herein to be highly expressed in cells that are either resistant or sensitive to the drug or compound. Alternatively, a patient's progress related to drug treatment or therapy can be monitored by obtaining a polynucleotide expression profile as described above, only after the patient has undergone treatment with a given drug or therapeutic compound. In this way, there is no need to test a patient sample prior to treatment with the drug or compound.

Such a monitoring process can indicate success or failure of a patient's treatment with a drug or compound based on the polynucleotide expression pattern of the cells isolated from the patient's sample, e.g., a tumor or cancer biopsy, as being relatively the same as or different from the polynucleotide expression pattern of the predictor polynucleotide set of the resistant or sensitive control panel of cells that have been exposed to the drug or compound and assessed for their polynucleotide expression profile following exposure. Thus, if, after treatment with a drug or compound, the test cells show a change in their polynucleotide expression profile from that seen prior to treatment to one which corresponds to that of the predictor polynucleotide set of the control panel of cells that are resistant to the drug or compound, it can serve as an indicator that the current treatment should be modified, changed, or even discontinued. Also, should a patient's response be one that shows sensitivity to treatment by a protein tyrosine kinase inhibitor compound, e.g., a Src tyrosine kinase inhibitor, based on correlation of the expression profile of the predictor polynucleotides of cells showing drug sensitivity with the polynucleotide expression profile from cells from a patient undergoing treatment, the patient's treatment prognosis can be qualified as favorable and treatment can continue. Further, if a patient has not been tested prior to drug treatment, the results obtained after treatment can be used to determine the resistance or sensitivity of the cells to the drug based on the polynucleotide expression profile compared with the predictor polynucleotide set.

In a related embodiment, the present invention embraces a method of monitoring the treatment of a patient having a disease treatable by a compound or agent that modulates a protein tyrosine kinase, i.e., breast cancer. Protein tyrosine kinases encompassed by such treatment monitoring assays include members of the Src

family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. For these assays, test cells from the patient are assayed to determine their polynucleotide expression pattern before and after exposure to a protein tyrosine kinase inhibitor compound or drug. The resulting polynucleotide expression profile of the cells tested before and after treatment is compared with the polynucleotide expression pattern of the predictor set of polynucleotides that have been described and shown herein to be highly expressed in cells that are either resistant or sensitive to the drug or compound. Thus, if a patient's response is or becomes one that is sensitive to treatment by a protein tyrosine kinase inhibitor compound, based on correlation of the expression profile of the predictor polynucleotides, the patient's treatment prognosis can be qualified as favorable and treatment can continue. Also, if after treatment with a drug or compound, the test cells do not exhibit a change in their polynucleotide expression profile to a profile that corresponds to that of the control panel of cells that are sensitive to the drug or compound, this serves as an indicator that the current treatment should be modified, changed, or even discontinued. Such monitoring processes can be repeated as necessary or desired and can indicate success or failure of a patient's treatment with a drug or compound, based on the polynucleotide expression pattern of the cells isolated from the patient's sample. The monitoring of a patient's response to a given drug treatment can also involve testing the patient's cells in the assay as described, only after treatment, rather than before and after treatment, with drug or active compound.

In a preferred embodiment, the present invention embraces a method of monitoring the treatment of a patient having a disease treatable by a compound or agent that modulates a src tyrosine kinase, i.e., breast cancer. The test cells from the patient are assayed to determine their polynucleotide expression pattern before and after exposure to a src tyrosine kinase inhibitor compound or drug. The resulting polynucleotide expression profile of the cells tested before and after treatment is compared with the polynucleotide expression pattern of the predictor set of polynucleotides that have been described and shown herein to be highly expressed in cells that are either resistant or sensitive to the drug or compound. Thus, if a patient's response is or becomes one that is sensitive to treatment by a src tyrosine kinase

inhibitor compound, based on correlation of the expression profile of the predictor polynucleotides, the patient's treatment prognosis can be qualified as favorable and treatment can continue. Also, if after treatment with a drug or compound, the test cells do not exhibit a change in their polynucleotide expression profile to a profile that
5 corresponds to that of the control panel of cells that are sensitive to the drug or compound, this serves as an indicator that the current treatment should be modified, changed, or even discontinued. Such monitoring processes can be repeated as necessary or desired and can indicate success or failure of a patient's treatment with a drug or compound, based on the polynucleotide expression pattern of the cells isolated
10 from the patient's sample. The monitoring of a patient's response to a given drug treatment can also involve testing the patient's cells in the assay as described only after treatment, rather than before and after treatment, with drug or active compound.

In another embodiment, the present invention encompasses a method of classifying whether a biological system, preferably cells from a tissue, organ, tumor or
15 cancer of an afflicted individual, will be resistant or sensitive to a compound that modulates the system. In a preferred aspect of this invention, the sensitivity or resistance of cells, e.g., those obtained from a tumor or cancer, to a protein tyrosine kinase inhibitor compound, or series of compounds, e.g., a Src tyrosine kinase inhibitor, is determined. Inhibitors can include those compounds, drugs, or biological
20 agents that inhibit, either directly or indirectly, the protein tyrosine kinases as described previously hereinabove. According to the method, a resistance/sensitivity profile of the cells after exposure to the protein tyrosine kinase inhibitor drug or compound can be determined via polynucleotide expression profiling protocols set forth herein. Such resistance/sensitivity profile of the cells reflects an IC_{50} value of
25 the cells to the compound(s) as determined using a suitable assay, such as an *in vitro* cytotoxicity assay as described in Example 1. A procedure of this sort can be performed using a variety of cell types and compounds that interact with the protein tyrosine kinase, or affect its activity in the signaling pathway of the protein tyrosine kinase.

30 In another of its embodiments, the present invention includes the preparation of one or more specialized microarrays (e.g., oligonucleotide microarrays or cDNA microarrays) comprising all of the polynucleotides in Tables 2, 4, or 5, or

combinations thereof, of the predictor polynucleotide sets described herein that have been demonstrated to be most highly correlated with sensitivity (or resistance) to protein tyrosine kinase modulators, particularly inhibitors of src tyrosine kinase. Preferably, the predictor polynucleotide sets are common for predicting sensitivity
5 among more than one protein tyrosine kinase modulator, e.g. a protein tyrosine kinase inhibitor such as a Src tyrosine kinase inhibitor, as demonstrated herein. In accordance with this aspect of the invention, the oligonucleotide sequences or cDNA sequences include any of the predictor polynucleotides or polynucleotide combinations as described herein, which are highly expressed in resistant or sensitive
10 cells, and are contained on a microarray, e.g., a oligonucleotide microarray or cDNA microarray in association with, or introduced onto, any supporting material, such as glass slides, nylon membrane filters, glass or polymer beads, chips, plates, or other types of suitable substrate material.

Cellular nucleic acid, e.g., RNA, is isolated either from cells undergoing
15 testing after exposure to a drug or compound that interacts with a protein tyrosine kinase as described herein, or its signaling pathway, or from cells being tested to obtain an initial determination or prediction of the cells' sensitivity to the drug or compound, and, ultimately, a prediction of treatment outcome with the drug or compound. The isolated nucleic acid is appropriately labeled and applied to one or
20 more of the specialized microarrays. The resulting pattern of polynucleotide expression on the specialized microarray is analyzed as described herein and known in the art. A pattern of polynucleotide expression correlating with either sensitivity or resistance to the drug or compound is able to be determined, e.g., via comparison with the polynucleotide expression pattern as shown in Figure 1 for the panel of cells
25 exposed to the protein tyrosine kinase inhibitor assayed herein.

In accordance with the specialized microarray embodiment of this invention, the microarray contains the polynucleotides of one or more of the predictor polynucleotide set(s) or subset(s), or a combination thereof, or all of the polynucleotides in Tables 2, 4, or 5, that are highly correlated with drug sensitivity or
30 resistance by a breast cell type. If the nucleic acid target isolated from test cells, such as tumor or cancer cells, preferably breast cancer or tumor cells, shows a high level of detectable binding to the polynucleotides of the predictor set for drug sensitivity

relative to control, then it can be predicted that a patient's cells will respond to the drug, or a series of drugs, and that the patient's response to the drug, or a series of drugs, will be favorable.

Such a result predicts that the cells of a tumor or cancer are good candidates
5 for the successful treatment or therapy utilizing the drug, or series of drugs. Alternatively, if the nucleic acid target isolated from test cells shows a high level of detectable binding to the polynucleotides of the predictor set for drug resistance, relative to control, then it can be predicted that a patient is likely not to respond to the drug, or a series of drugs, and that the patient's response to the drug, or a series of
10 drugs, is not likely to be favorable. Such a result predicts that the cells of a tumor or cancer are not good candidates for treatment or therapy utilizing the drug, or series of drugs.

The utilization of microarray technology is known and practiced in the art. Briefly, to determine polynucleotide expression using microarray technology,
15 polynucleotides, e.g., RNA, DNA, cDNA, preferably RNA, are isolated from a biological sample, e.g., cells, as described herein for breast cells, using procedures and techniques that are practiced in the art. The isolated nucleic acid is detectably labeled, e.g., fluorescent, enzyme, radionuclide, or chemiluminescent label, and applied to a microarray, e.g., the specialized microarrays provided by this invention. The array is
20 then washed to remove unbound material and visualized by staining or fluorescence, or other means known in the art depending on the type of label utilized.

In another embodiment of this invention, the predictor polynucleotides (Table 2), or one or more subsets of polynucleotides comprising the predictor polynucleotide sets (e.g., Tables 4-5) can be used as biomarkers for cells that are resistant or sensitive
25 to protein tyrosine kinase inhibitor compounds, e.g., Src tyrosine kinase inhibitors. With the predictor polynucleotides in hand, screening and detection assays can be carried out to determine whether or not a given compound, preferably a protein tyrosine kinase inhibitor compound such as a Src tyrosine kinase inhibitor compound, elicits a sensitive or a resistant phenotype following exposure of cells, e.g., cells taken
30 from a tumor or cancer biopsy sample, such as a breast cancer cell sample, to the compound. Thus, methods of screening, monitoring, detecting, prognosing and/or diagnosing to determine the resistance or sensitivity of cells to a drug or compound

that interacts with a protein tyrosine kinase, or a protein tyrosine kinase pathway, preferably an inhibitor compound, and to which the cells are exposed, are encompassed by the present invention.

Such methods embrace a variety of procedures and assays to determine and
5 assess the expression of polynucleotides, in particular, the predictor or src biomarker polynucleotides and predictor polynucleotide subsets as described herein (Tables 2, 4, and 5), in cells that have been exposed to drugs or compounds that interact with or effect a protein tyrosine kinase, or a protein tyrosine kinase pathway, wherein the protein tyrosine kinases include members of the Src family of tyrosine kinases, for
10 example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Suitable methods include detection and evaluation of polynucleotide activation or expression at the level of nucleic acid, e.g., DNA, RNA, mRNA, and detection and evaluation of encoded protein. For example, PCR assays as known and practiced in the art can be
15 employed to quantify RNA or DNA in cells being assayed for susceptibility to drug treatment, for example, protein tyrosine kinase inhibitors. (see Example 2, RT-PCR).

In another embodiment, the present invention is directed to a method of identifying cells, tissues, and/or patients that are predicted to be resistant to either protein tyrosine inhibitor compounds or compounds that affect protein tyrosine kinase
20 signaling pathways, e.g., Src tyrosine kinase, or that are resistant in different biological systems to those compounds. The method comprises the step(s) of (i) analyzing the expression of only those polynucleotides listed in Tables 2, 4, 5, or any combination thereof, that have been shown to be correlative to predicting resistant responses to such compounds; (ii) comparing the observed expression levels of those
25 correlative resistant polynucleotides in the test cells, tissues, and/or patients to the expression levels of those same polynucleotides in a cell line that is known to be resistant to the compounds; and (iii) predicting whether the cells, tissues, and/or patients are resistant to the compounds based upon the overall similarity of the observed expression of those polynucleotides in step (ii).

30 In another embodiment, the present invention is directed to a method of identifying cells, tissues, and/or patients that are predicted to be sensitive to either protein tyrosine inhibitor compounds or compounds that affect protein tyrosine kinase

signaling pathways, e.g., the Src tyrosine kinase, or that are sensitive in different biological systems to those compounds. The method involves the step(s) of (i) analyzing the expression of only those polynucleotides listed in Tables 2, 4, 5, or any combination thereof, that have been shown to be correlative to predicting sensitive
5 responses to such compounds; (ii) comparing the observed expression levels of those correlative sensitive polynucleotides in the test cells, tissues, and/or patients to the expression levels of those same polynucleotides in a cell line that is known to be sensitive to the compounds; and (iii) predicting whether the cells, tissues, and/or patients are sensitive to the compounds based upon the overall similarity of the
10 observed expression of those polynucleotides in step (ii).

The present invention further encompasses the detection and/or quantification of one or more of the protein tyrosine kinase biomarker proteins of the present invention using antibody-based assays (e.g., immunoassays) and/or detection systems. As mentioned, protein tyrosine kinases encompass members of the Src family of
15 tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Such assays include the following non-limiting examples, ELISA, immunofluorescence, fluorescence activated cell sorting (FACS), Western Blots, etc., as further described herein.

20 In another embodiment, the human protein tyrosine kinase biomarker polypeptides and/or peptides of the present invention, or immunogenic fragments or oligopeptides thereof, can be used for screening therapeutic drugs or compounds in a variety of drug screening techniques. The fragment employed in such a screening assay can be free in solution, affixed to a solid support, borne on a cell surface, or
25 located intracellularly. The reduction or abolition of activity of the formation of binding complexes between the biomarker protein and the agent being tested can be measured. Thus, the present invention provides a method for screening or assessing a plurality of compounds for their specific binding affinity with a protein kinase inhibitor biomarker polypeptide, or a bindable peptide fragment thereof, of this
30 invention. The method comprises the steps of providing a plurality of compounds; combining the protein kinase inhibitor biomarker polypeptide, or a bindable peptide fragment thereof, with each of the plurality of compounds, for a time sufficient to

allow binding under suitable conditions; and detecting binding of the biomarker polypeptide or peptide to each of the plurality of test compounds, thereby identifying the compounds that specifically bind to the biomarker polypeptide or peptide. More specifically, the biomarker polypeptide or peptide is that of a Src tyrosine kinase inhibitor biomarkers.

Methods to identify compounds that modulate the activity of the human protein tyrosine kinase biomarker polypeptides and/or peptides provided in Table 2 by the present invention, comprise combining a candidate compound or drug modulator of protein kinases and measuring an effect of the candidate compound or drug modulator on the biological activity of the protein kinase inhibitor biomarker polypeptide or peptide. Such measurable effects include, for example, a physical binding interaction; the ability to cleave a suitable protein kinase substrate; effects on a native and cloned protein kinase biomarker-expressing cell line; and effects of modulators or other protein kinase-mediated physiological measures.

Another method of identifying compounds that modulate the biological activity of the protein tyrosine kinase biomarker polypeptides of the present invention comprises combining a potential or candidate compound or drug modulator of a protein tyrosine kinase biological activity, e.g., a Src tyrosine kinase, with a host cell that expresses the protein tyrosine kinase biomarker polypeptide and measuring an effect of the candidate compound or drug modulator on the biological activity of the protein tyrosine kinase biomarker polypeptides. The host cell can also be capable of being induced to express the protein tyrosine kinase biomarker polypeptide, e.g., via inducible expression. Physiological effects of a given modulator candidate on the protein tyrosine kinase biomarker polypeptide can also be measured. Thus, cellular assays for particular protein tyrosine kinase modulators, e.g., a src kinase modulator, can be either direct measurement or quantification of the physical biological activity of the protein tyrosine kinase biomarker polypeptide, or they may be measurement or quantification of a physiological effect. Such methods preferably employ a protein tyrosine kinase biomarker polypeptide as described herein, or an overexpressed recombinant protein tyrosine kinase biomarker polypeptide in suitable host cells containing an expression vector as described herein, wherein the protein tyrosine

kinase biomarker polypeptide is expressed, overexpressed, or undergoes up-regulated expression.

Another aspect of the present invention embraces a method of screening for a compound that is capable of modulating the biological activity of a protein tyrosine kinase biomarker polypeptide, e.g., a Src kinase biomarker polypeptide. The method comprises providing a host cell containing an expression vector harboring a nucleic acid sequence encoding a protein tyrosine kinase biomarker polypeptide, or a functional peptide or portion thereof (e.g., the src polypeptide, protein, peptide, or fragment sequences as set forth in Table 2, or the Sequence Listing herein); determining the biological activity of the expressed protein tyrosine kinase biomarker polypeptide in the absence of a modulator compound; contacting the cell with the modulator compound and determining the biological activity of the expressed protein tyrosine kinase biomarker polypeptide in the presence of the modulator compound. In such a method, a difference between the activity of the protein tyrosine kinase biomarker polypeptide in the presence of the modulator compound and in the absence of the modulator compound indicates a modulating effect of the compound.

Essentially any chemical compound can be employed as a potential modulator or ligand in the assays according to the present invention. Compounds tested as protein tyrosine kinase modulators can be any small chemical compound, or biological entity (e.g., protein, sugar, nucleic acid, or lipid). Test compounds are typically small chemical molecules and peptides. Generally, the compounds used as potential modulators can be dissolved in aqueous or organic (e.g., DMSO-based) solutions. The assays are designed to screen large chemical libraries by automating the assay steps and providing compounds from any convenient source. Assays are typically run in parallel, for example, in microtiter formats on microtiter plates in robotic assays. There are many suppliers of chemical compounds, including, for example, Sigma (St. Louis, MO), Aldrich (St. Louis, MO), Sigma-Aldrich (St. Louis, MO), Fluka Chemika-Biochemica Analytika (Buchs, Switzerland). Also, compounds can be synthesized by methods known in the art.

High throughput screening methodologies are particularly envisioned for the detection of modulators of the novel protein tyrosine kinase biomarker, e.g., src biomarker, polynucleotides and polypeptides described herein. Such high throughput

screening methods typically involve providing a combinatorial chemical or peptide library containing a large number of potential therapeutic compounds (e.g., ligand or modulator compounds). The combinatorial chemical libraries or ligand libraries are then screened in one or more assays to identify those library members (e.g., particular
5 chemical species or subclasses) that display a desired characteristic activity. The compounds so identified can serve as conventional lead compounds, or can themselves be used as potential or actual therapeutics.

A combinatorial chemical library is a collection of diverse chemical compounds generated either by chemical synthesis or biological synthesis, prepared
10 by combining a number of chemical building blocks (i.e., reagents such as amino acids). As an example, a linear combinatorial library, e.g., a polypeptide or peptide library, is formed by combining a set of chemical building blocks in every possible way for a given compound length (i.e., the number of amino acids in a polypeptide or peptide compound). Millions of chemical compounds can be synthesized through
15 such combinatorial mixing of chemical building blocks.

The preparation and screening of combinatorial chemical libraries is well known to those having skill in the pertinent art. Combinatorial libraries include, without limitation, peptide libraries (e.g. U.S. Patent No. 5,010,175; Furka, 1991, *Int. J. Pept. Prot. Res.*, 37:487-493; and Houghton et al., 1991, *Nature*, 354:84-88). Other
20 chemistries for generating chemical diversity libraries can also be used. Nonlimiting examples of chemical diversity library chemistries include, peptoids (PCT Publication No. WO 91/019735), encoded peptides (PCT Publication No. WO 93/20242), random bio-oligomers (PCT Publication No. WO 92/00091), benzodiazepines (U.S. Patent No. 5,288,514), diversomers such as hydantoins, benzodiazepines and dipeptides
25 (Hobbs et al., 1993, *Proc. Natl. Acad. Sci. USA*, 90:6909-6913), vinylogous polypeptides (Hagihara et al., 1992, *J. Amer. Chem. Soc.*, 114:6568), nonpeptidal peptidomimetics with glucose scaffolding (Hirschmann et al., 1992, *J. Amer. Chem. Soc.*, 114:9217-9218), analogous organic synthesis of small compound libraries (Chen et al., 1994, *J. Amer. Chem. Soc.*, 116:2661), oligocarbamates (Cho et al., 1993, *Science*, 261:1303), and/or peptidyl phosphonates (Campbell et al., 1994, *J. Org. Chem.*, 59:658), nucleic acid libraries (see Ausubel, Berger and Sambrook, all supra),
30 peptide nucleic acid libraries (U.S. Patent No. 5,539,083), antibody libraries (e.g.,

Vaughn et al., 1996, *Nature Biotechnology*, 14(3):309-314) and PCT/US96/10287), carbohydrate libraries (e.g., Liang et al., 1996, *Science*, 274-1520-1522) and U.S. Patent No. 5,593,853), small organic molecule libraries (e.g., benzodiazepines, Baum C&EN, Jan. 18, 1993, page 33; and U.S. Patent No. 5,288,514; isoprenoids (U.S. Patent No. 5,569,588); thiazolidinones and metathiazanones (U.S. Patent No. 5,549,974); pyrrolidines (U.S. Patent Nos. 5,525,735 and 5,519,134); morpholino compounds (U.S. Patent No. 5,506,337); and the like.

Devices for the preparation of combinatorial libraries are commercially available (e.g., 357 MPS, 390 MPS, Advanced Chem Tech, Louisville KY; Symphony, Rainin, Woburn, MA; 433A Applied Biosystems, Foster City, CA; 9050 Plus, Millipore, Bedford, MA). In addition, a large number of combinatorial libraries are commercially available (e.g., ComGenex, Princeton, NJ; Asinex, Moscow, Russia; Tripos, Inc., St. Louis, MO; ChemStar, Ltd., Moscow, Russia; 3D Pharmaceuticals, Exton, PA; Martek Biosciences, Columbia, MD, and the like).

In one aspect, the invention provides solid phase-based *in vitro* assays in a high throughput format, where the cell or tissue expressing a tyrosine kinase protein/polypeptide/peptide is attached to a solid phase substrate. In such high throughput assays, it is possible to screen up to several thousand different modulators or ligands in a single day. In particular, each well of a microtiter plate can be used to perform a separate assay against a selected potential modulator, or, if concentration or incubation time effects are to be observed, every 5-10 wells can be used to test a single modulator. Thus, a single standard microtiter plate can be used in to assay about 96 modulators. If 1536 well plates are used, then a single plate can easily assay from about 100 to about 1500 different compounds. It is possible to assay several different plates per day; thus, for example, assay screens for up to about 6,000-20,000 different compounds are possible using the described integrated systems.

In another of its aspects, the present invention encompasses screening and small molecule (e.g., drug) detection assays which involve the detection or identification of small molecules that can bind to a given protein, i.e., a tyrosine kinase biomarker polypeptide or peptide, such as a Src tyrosine kinase biomarker polypeptide or peptide. Particularly preferred are assays suitable for high throughput screening methodologies.

In such binding-based detection, identification, or screening assays, a functional assay is not typically required. All that is needed, in general, is a target protein, preferably substantially purified, and a library or panel of compounds (e.g., ligands, drugs, or small molecules), or biological entities to be screened or assayed for
5 binding to the protein target. Preferably, most small molecules that bind to the target protein modulate the target's activity in some manner due to preferential, higher affinity binding to functional areas or sites on the protein.

An example of such an assay is the fluorescence based thermal shift assay (3-Dimensional Pharmaceuticals, Inc., 3DP, Exton, PA) as described in U.S. Patent Nos.
10 6,020,141 and 6,036,920 to Pantoliano et al. (See also, J. Zimmerman, 2000, *Gen. Eng. News*, 20(8)). The assay allows the detection of small molecules (e.g., drugs, ligands) that bind to expressed, and preferably purified, tyrosine kinase biomarker proteins/polypeptides/peptides, such as the Src tyrosine kinase, based on affinity of binding determinations by analyzing thermal unfolding curves of protein-drug or
15 ligand complexes. The drugs or binding molecules determined by this technique can be further assayed, if desired, by methods such as those described herein to determine if the molecules affect or modulate function or activity of the target protein.

To purify a tyrosine kinase biomarker polypeptide or peptide, e.g., Src tyrosine kinase, to measure a biological binding or ligand binding activity, the source may be a
20 whole cell lysate that can be prepared by successive freeze-thaw cycles (e.g., one to three) in the presence of standard protease inhibitors. The tyrosine kinase biomarker polypeptide can be partially or completely purified by standard protein purification methods, e.g., affinity chromatography using specific antibody(ies) described herein, or by ligands specific for an epitope tag engineered into the recombinant tyrosine
25 kinase biomarker polypeptide molecule, also as described herein. Binding activity can then be measured as described.

Compounds which are identified according to the methods provided herein, and which modulate or regulate the biological activity or physiology of the tyrosine kinase biomarker polypeptides according to the present invention, are a preferred
30 embodiment of this invention. It is contemplated that such modulatory compounds can be employed in treatment and therapeutic methods for treating a condition that is mediated by the tyrosine kinase biomarker polypeptides, e.g., Src tyrosine kinase

biomarker polypeptides, by administering to an individual in need of such treatment a therapeutically effective amount of the compound identified by the methods described herein.

In addition, the present invention provides methods for treating an individual
5 in need of such treatment for a disease, disorder, or condition that is mediated by the tyrosine kinase biomarker polypeptides of the invention, comprising administering to the individual a therapeutically effective amount of the tyrosine kinase biomarker-modulating compound identified by a method provided herein. In accordance with this invention, the tyrosine kinase biomarker polypeptides or proteins encompassed by
10 the methods include members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors.

The present invention particularly provides methods for treating an individual in need of such treatment for a disease, disorder, or condition that is mediated by Src
15 biomarker polypeptides of the invention, comprising administering to the individual a therapeutically effective amount of the Src biomarker-modulating compound identified by a method provided herein.

The present invention further encompasses polypeptides comprising, or alternatively consisting of, an epitope of the polypeptide having an amino acid
20 sequence of one or more of the protein tyrosine kinase biomarkers, preferably the Src biomarker amino acid sequences as set forth in Table 2. The present invention also encompasses polynucleotide sequences encoding an epitope of a polypeptide sequence of the protein tyrosine kinase biomarkers of the invention.

The term "epitopes" as used herein, refers to portions of a polypeptide having
25 antigenic or immunogenic activity in an animal, preferably a mammal, and most preferably a human. In a preferred embodiment, the present invention encompasses a polypeptide comprising an epitope, as well as the polynucleotide encoding this polypeptide. An "immunogenic epitope" as used herein, refers to a portion of a protein that elicits an antibody response in an animal, as determined by any method
30 known in the art, for example, by the methods for generating antibodies described *infra*. (See, for example, Geysen et al., 1983, *Proc. Natl. Acad. Sci. USA*, 81:3998-4002). The term "antigenic epitope" as used herein refers to a portion of a protein to

which an antibody can immunospecifically bind to its antigen as determined by any method well known in the art, for example, by the immunoassays described herein. Immunospecific binding excludes non-specific binding, but does not necessarily exclude cross-reactivity with other antigens. Antigenic epitopes need not necessarily be immunogenic. Either the full-length protein or an antigenic peptide fragment can be used. Antibodies are preferably prepared from these regions or from discrete fragments in regions of the tyrosine kinase biomarker nucleic acid and protein sequences comprising an epitope. Polypeptide or peptide fragments that function as epitopes may be produced by any conventional means. (See, e.g., Houghten, 1985, *Proc. Natl. Acad. Sci. USA*, 82:5131-5135; and as described in U. S. Patent No. 4,631,211).

Moreover, antibodies can also be prepared from any region of the polypeptides and peptides of the protein tyrosine kinase biomarkers, including Src kinase biomarkers as described herein. In addition, if a polypeptide is a receptor protein, antibodies can be developed against an entire receptor or portions of the receptor, for example, the intracellular carboxy terminal domain, the amino terminal extracellular domain, the entire transmembrane domain, specific transmembrane segments, any of the intracellular or extracellular loops, or any portions of these regions. Antibodies can also be developed against specific functional sites, such as the site of ligand binding, or sites that are glycosylated, phosphorylated, myristylated, or amidated, for example.

In the present invention, antigenic epitopes for generating antibodies preferably contain a sequence of at least 4, at least 5, at least 6, at least 7, more preferably at least 8, at least 9, at least 10, at least 11, at least 12, at least 13, at least 14, at least 15, at least 20, at least 25, at least 30, at least 40, at least 50, and, most preferably, between about 15 to about 30 amino acid residues. Combinations of the foregoing epitopes are included. Preferred polypeptides comprising immunogenic or antigenic epitopes are at least 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, or 100 amino acid residues in length. Additional non-exclusive preferred antigenic epitopes include the antigenic epitopes disclosed herein, as well as portions thereof, as well as any combination of two, three, four, five or more of these antigenic epitopes. Such antigenic epitopes can be used as the target molecules in

immunoassays. (See, for instance, Wilson et al., 1984, *Cell*, 37:767-778; and Sutcliffe et al., 1983, *Science*, 219:660-666). The fragments as described herein are not to be construed, however, as encompassing any fragments which may be disclosed prior to the invention.

5 Protein tyrosine kinase biomarker polypeptides comprising one or more immunogenic epitopes which elicit an antibody response can be introduced together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse). Alternatively, if the polypeptide is of sufficient length (e.g., at least about 15-25 amino acids), the polypeptide can be presented without a carrier. However,
10 immunogenic epitopes comprising as few as 5 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting).

Epitope-bearing polypeptides of the present invention can be used to induce antibodies according to methods well known in the art including, but not limited to, *in vivo* immunization, *in vitro* immunization, and phage display methods. See, e. g.,
15 Sutcliffe et al., *supra*; Wilson et al., *supra*; and Bittle et al., *supra*). If *in vivo* immunization is used, animals can be immunized with free peptide of appropriate size; however, the anti-peptide antibody titer can be boosted by coupling the peptide to a macromolecular carrier, such as keyhole limpet hemacyanin (KLH), or tetanus
20 toxoid (TT). For instance, peptides containing cysteine residues can be coupled to a carrier using a linker such as maleimidobenzoyl-N-hydroxysuccinimide ester (MBS), while other peptides may be coupled to carriers using a more general linking agent, such as glutaraldehyde.

Peptides containing epitopes can also be synthesized as multiple antigen
25 peptides (MAPs), first described by J.P. Tam et al. (1995, *Biomed. Pept., Proteins, Nucleic Acids*, 199, 1(3):123-32) and Calvo et al. (1993, *J. Immunol.*, 150(4):1403-12), which are hereby incorporated by reference in their entirety herein. MAPs contain multiple copies of a specific peptide attached to a non-immunogenic lysine core. MAP peptides usually contain four or eight copies of the peptide, which are
30 often referred to as MAP4 or MAP8 peptides. By way of non-limiting example, MAPs can be synthesized onto a lysine core matrix attached to a polyethylene glycol-polystyrene (PEG-PS) support. The peptide of interest is synthesized onto the lysine

residues using 9-fluorenylmethoxycarbonyl (Fmoc) chemistry. For example, Applied Biosystems (Foster City, CA) offers commercially available MAP resins, such as, for example, the Fmoc Resin 4 Branch and the Fmoc Resin 8 Branch which can be used to synthesize MAPs. Cleavage of MAPs from the resin is performed with standard
5 trifluoroacetic acid (TFA)-based cocktails known in the art. Purification of MAPs, except for desalting, is not generally necessary. MAP peptides can be used in immunizing vaccines which elicit antibodies that recognize both the MAP and the native protein from which the peptide was derived.

Epitope-bearing peptides of the invention can also be incorporated into a coat
10 protein of a virus, which can then be used as an immunogen or a vaccine with which to immunize animals, including humans, in order stimulate the production of anti-epitope antibodies. For example, the V3 loop of the gp120 glycoprotein of the human immunodeficiency virus type 1 (HIV-1) has been engineered to be expressed on the surface of rhinovirus. Immunization with rhinovirus displaying the V3 loop peptide
15 yielded apparently effective mimics of the HIV-1 immunogens (as measured by their ability to be neutralized by anti-HIV-1 antibodies as well as by their ability to elicit the production of antibodies capable of neutralizing HIV-1 in cell culture). This techniques of using engineered viral particles as immunogens is described in more detail in Smith et al., 1997, *Behring Inst Mitt Feb*, (98):229-39; Smith et al., 1998, *J.*
20 *Virol.*, 72:651-659; and Zhang et al., 1999, *Biol. Chem.*, 380:365-74), which are hereby incorporated by reference herein in their entireties.

Moreover, polypeptides or peptides containing epitopes according to the present invention can be modified, for example, by the addition of amino acids at the amino- and/or carboxy-terminus of the peptide. Such modifications are performed,
25 for example, to alter the conformation of the epitope bearing polypeptide such that the epitope will have a conformation more closely related to the structure of the epitope in the native protein. An example of a modified epitope-bearing polypeptide of the invention is a polypeptide in which one or more cysteine residues have been added to the polypeptide to allow for the formation of a disulfide bond between two cysteines,
30 thus resulting in a stable loop structure of the epitope-bearing polypeptide under non-reducing conditions. Disulfide bonds can form between a cysteine residue added to the polypeptide and a cysteine residue of the naturally-occurring epitope, or between

two cysteines which have both been added to the naturally-occurring epitope-bearing polypeptide.

In addition, it is possible to modify one or more amino acid residues of the naturally-occurring epitope-bearing polypeptide by substitution with cysteines to promote the formation of disulfide bonded loop structures. Cyclic thioether molecules of synthetic peptides can be routinely generated using techniques known in the art, e.g., as described in PCT publication WO 97/46251, incorporated in its entirety by reference herein. Other modifications of epitope-bearing polypeptides contemplated by this invention include biotinylation.

For the production of antibodies *in vivo*, host animals, such as rabbits, rats, mice, sheep, or goats, are immunized with either free or carrier-coupled peptides or MAP peptides, for example, by intraperitoneal and/or intradermal injection. Injection material is typically an emulsion containing about 100 µg of peptide or carrier protein and Freund's adjuvant, or any other adjuvant known for stimulating an immune response. Several booster injections may be needed, for instance, at intervals of about two weeks, to provide a useful titer of anti-peptide antibody which can be detected, for example, by ELISA assay using free peptide adsorbed to a solid surface. The titer of anti-peptide antibodies in serum from an immunized animal can be increased by selection of anti-peptide antibodies, e.g., by adsorption of the peptide onto a solid support and elution of the selected antibodies according to methods well known in the art.

As one having skill in the art will appreciate, and as discussed above, the tyrosine kinase biomarker polypeptides of the present invention, which include the following: e.g., members of the Src family of tyrosine kinases, such as Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors, and which comprise an immunogenic or antigenic epitope, can be fused to other polypeptide sequences. For example, the polypeptides of the present invention can be fused with the constant domain of immunoglobulins (IgA, IgE, IgG, IgD, or IgM), or portions thereof, e.g., CH1, CH2, CH3, or any combination thereof, and portions thereof, or with albumin (including, but not limited to, recombinant human albumin, or fragments or variants thereof (see, e. g., U. S. Patent No. 5,876,969; EP Patent No. 0 413 622; and U.S. Patent No.

5,766,883, incorporated by reference in their entirety herein), thereby resulting in chimeric polypeptides. Such fusion proteins may facilitate purification and may increase half-life *in vivo*. This has been shown for chimeric proteins containing the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (see, e.g.,
5 Traunecker et al., 1988, *Nature*, 331:84-86).

Enhanced delivery of an antigen across the epithelial barrier to the immune system has been demonstrated for antigens (e.g., insulin) conjugated to an FcRn binding partner, such as IgG or Fc fragments (see, e.g., WO 96/22024 and WO
10 99/04813). IgG fusion proteins that have a disulfide-linked dimeric structure due to the IgG portion disulfide bonds have also been found to be more efficient in binding and neutralizing other molecules than are monomeric polypeptides, or fragments thereof, alone. (See, e.g., Fountoulakis et al., 1995, *J. Biochem.*, 270:3958-3964).

Nucleic acids encoding epitopes can also be recombined with a polynucleotide of interest as an epitope tag (e.g., the hemagglutinin ("HA") tag or flag tag) to aid in
15 detection and purification of the expressed polypeptide. For example, a system for the ready purification of non-denatured fusion proteins expressed in human cell lines has been described by Janknecht et al., (1991, *Proc. Natl. Acad. Sci. USA*, 88:8972- 897). In this system, the polynucleotide of interest is subcloned into a vaccinia recombination plasmid such that the open reading frame of the polynucleotide is
20 translationally fused to an amino-terminal tag having six histidine residues. The tag serves as a matrix binding domain for the fusion protein. Extracts from cells infected with the recombinant vaccinia virus are loaded onto an Ni²⁺ nitriloacetic acid-agarose column and histidine-tagged proteins are selectively eluted with imidazole-containing
25 buffers.

Additional fusion proteins of the invention can be generated by employing the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling"). DNA shuffling can be employed to modulate the activities of polypeptides of the invention; such methods can be used to
30 generate polypeptides with altered activity, as well as agonists and antagonists of the polypeptides. See, generally, U.S. Patent Nos. 5,605,793; 5,811,238; 5,830,721; 5,834,252; and 5,837,458, and Patten et al., 1997, *Curr. Opinion Biotechnol.*, 8:724-

33; Harayama, 1998, *Trends Biotechnol.*, 16(2):76-82; Hansson, et al., 1999, *J. Mol. Biol.*, 287:265-76; and Lorenzo and Blasco, 1998, *Biotechniques*, 24(2):308-313, the contents of each of which are hereby incorporated by reference in its entirety.

In an embodiment of the invention, alteration of polynucleotides
5 corresponding to one or more of the src biomarker polynucleotide sequences as set forth in Table 2, and the polypeptides encoded by these polynucleotides, can be achieved by DNA shuffling. DNA shuffling involves the assembly of two or more DNA segments by homologous or site-specific recombination to generate variation in the polynucleotide sequence. In another embodiment, polynucleotides of the
10 invention, or their encoded polypeptides, may be altered by being subjected to random mutapolynucleotidesis by error-prone PCR, random nucleotide insertion, or other methods, prior to recombination. In another embodiment, one or more components, motifs, sections, parts, domains, fragments, etc., of a polynucleotide encoding a polypeptide of this invention may be recombined with one or more components,
15 motifs, sections, parts, domains, fragments, etc. of one or more heterologous molecules.

Another aspect of the present invention relates to antibodies and T-cell antigen receptors (TCRs), which immunospecifically bind to a polypeptide, polypeptide fragment, or variant one or more of the src biomarker amino acid sequences as set
20 forth in Table 2, and/or an epitope thereof, of the present invention (as determined by immunoassays well known in the art for assaying specific antibody-antigen binding).

A bispecific or bifunctional antibody is an artificial hybrid antibody having two different heavy/light chain pairs and two different binding sites. Bispecific antibodies can be produced by a variety of methods, including fusion of hybridomas
25 or linking of Fab' fragments. (See, e. g., Songsivilai & Lachmann, 1990, *Clin. Exp. Immunol.*, 79:315-321; Kostelny et al., 1992, *J. Immunol.*, 148:1547-1553). In addition, bispecific antibodies can be formed as "diabodies" (See, Holliger et al., 1993, *Proc. Natl. Acad. Sci. USA*, 90:6444-6448), or "Janusins" (See, Traunecker et al., 1991, *EMBO J.*, 10:3655-3659 and Traunecker et al., 1992, *Int. J. Cancer Suppl.*
30 7:51-52-127).

Antibodies of the invention include, but are not limited to, polyclonal, monoclonal, multispecific, human, humanized or chimeric antibodies, single chain

antibodies, Fab fragments, F(ab') fragments, fragments produced by a Fab expression library, anti-idiotypic (anti-Id) antibodies (including, e.g., anti-Id antibodies to antibodies of the invention), intracellularly made antibodies (i.e., intrabodies), and epitope-binding fragments of any of the above. The term "antibody", as used herein, 5 refers to immunoglobulin molecules and immunologically active portions or fragments of immunoglobulin molecules, i.e., molecules that contain an antigen binding site that immunospecifically binds an antigen. The immunoglobulin molecules of the invention can be of any type (e.g., IgG, IgE, IgM, IgD, IgA and IgY), class or subclass (e.g., IgG1, IgG2, IgG3, IgG4, IgA1 and IgA2) of immunoglobulin 10 molecule. Preferably, immunoglobulin is an IgG1, an IgG2, or an IgG4 isotype.

Immunoglobulins may have both a heavy and a light chain. An array of IgG, IgE, IgM, IgD, IgA, and IgY heavy chains can be paired with a light chain of the kappa or lambda types. Most preferably, the antibodies of the present invention are human antigen-binding antibodies and antibody fragments and include, but are not 15 limited to, Fab, Fab' F(ab')₂, Fd, single-chain Fvs (scFv), single-chain antibodies, disulfide-linked Fvs (sdFv) and fragments comprising either a V_L or V_H domain. Antigen-binding antibody fragments, including single-chain antibodies, can comprise the variable region(s) alone or in combination with the entirety or a portion of the following: hinge region, and CH1, CH2, and CH3 domains. Also included in 20 connection with the invention are antigen-binding fragments comprising any combination of variable region(s) with a hinge region, and CH1, CH2, and CH3 domains. The antibodies of the invention can be from any animal origin including birds and mammals. Preferably, the antibodies are of human, murine (e. g., mouse and rat), donkey, sheep, rabbit, goat, guinea pig, camel, horse, or chicken origin. As 25 used herein, "human" antibodies include antibodies having the amino acid sequence of a human immunoglobulin and include antibodies isolated from human immunoglobulin libraries or from animals transgenic for one or more human immunoglobulin and that do not express endogenous immunoglobulins, as described *infra* and, for example, in U.S. Patent No. 5,939,598.

30 The antibodies of the present invention can be monospecific, bispecific, trispecific, or of greater multispecificity. Multispecific antibodies can be specific for different epitopes of a polypeptide of the present invention, or can be specific for both

a polypeptide of the present invention, and a heterologous epitope, such as a heterologous polypeptide or solid support material. (See, e.g., WO 93/17715; WO 92/08802; WO 91/00360; WO 92/05793; Tutt et al., 1991, *J. Immunol.*, 147:60-69; U.S. Patent Nos. 4,474,893; 4,714,681; 4,925,648; 5,573,920; 5,601,819; and
5 Kostelny et al., 1992, *J. Immunol.*, 148:1547-1553).

Antibodies of the present invention can be described or specified in terms of the epitope(s) or portion(s) of a polypeptide of the present invention which they recognize or specifically bind. The epitope(s) or polypeptide portion(s) can be specified, e.g., by N-terminal and C-terminal positions, by size in contiguous amino
10 acid residues, or as presented in the sequences defined in Table 2 herein. Further included in accordance with the present invention are antibodies which bind to polypeptides encoded by polynucleotides which hybridize to a polynucleotide of the present invention under stringent, or moderately stringent, hybridization conditions as described herein.

15 The antibodies of the invention (including molecules comprising, or alternatively consisting of, antibody fragments or variants thereof) can bind immunospecifically to a polypeptide or polypeptide fragment or to a variant human protein tyrosine kinase biomarker of the invention, e.g., the Src biomarker proteins as set forth in Table 2, and/or monkey src biomarker protein.

20 By way of non-limiting example, an antibody can be considered to bind to a first antigen preferentially if it binds to the first antigen with a dissociation constant (Kd) that is less than the antibody's Kd for the second antigen. In another non-limiting embodiment, an antibody can be considered to bind to a first antigen preferentially if it binds to the first antigen with an affinity that is at least one order of
25 magnitude less than the antibody's Ka for the second antigen. In another non-limiting embodiment, an antibody can be considered to bind to a first antigen preferentially if it binds to the first antigen with an affinity that is at least two orders of magnitude less than the antibody's Kd for the second antigen.

In another nonlimiting embodiment, an antibody may be considered to bind to
30 a first antigen preferentially if it binds to the first antigen with an off rate (koff) that is less than the antibody's koff for the second antigen. In another nonlimiting embodiment, an antibody can be considered to bind to a first antigen preferentially if

it binds to the first antigen with an affinity that is at least one order of magnitude less than the antibody's koff for the second antigen. In another nonlimiting embodiment, an antibody can be considered to bind to a first antigen preferentially if it binds to the first antigen with an affinity that is at least two orders of magnitude less than the
5 antibody's koff for the second antigen.

Antibodies of the present invention can also be described or specified in terms of their binding affinity to a tyrosine kinase biomarker polypeptide of the present invention, e.g., members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including,
10 Bcr-abl, Jak, PDGFR, c-kit and Eph receptors. Preferred binding affinities include those with a dissociation constant or Kd of less than 5×10^{-2} M, 1×10^{-2} M, 5×10^{-3} M, 1×10^{-3} M, 5×10^{-4} M, or 1×10^{-4} M. More preferred binding affinities include those with a dissociation constant or Kd less than 5×10^{-5} M, 1×10^{-5} M, 5×10^{-6} M, 1×10^{-6} M, 5×10^{-7} M, 1×10^{-7} M, 5×10^{-8} M, or 1×10^{-8} M. Even more preferred
15 antibody binding affinities include those with a dissociation constant or Kd of less than 5×10^{-9} M, 1×10^{-9} M, 5×10^{-10} M, 1×10^{-10} M, 5×10^{-11} M, 1×10^{-11} M, 5×10^{-12} M, 1×10^{-12} M, 5×10^{-13} M, 1×10^{-13} M, 5×10^{-14} M, 1×10^{-14} M, 5×10^{-15} M, or 1×10^{-15} M.

In specific embodiments, antibodies of the invention bind to the protein
20 tyrosine kinase biomarker polypeptides, or fragments, or variants thereof, with an off rate (koff) of less than or equal to about $5 \times 10^{-2} \text{ sec}^{-1}$, $1 \times 10^{-2} \text{ sec}^{-1}$, $5 \times 10^{-3} \text{ sec}^{-1}$, or $1 \times 10^{-3} \text{ sec}^{-1}$. More preferably, antibodies of the invention bind to src biomarker protein polypeptides or fragments or variants thereof with an off rate (koff) of less than or equal to about $5 \times 10^{-4} \text{ sec}^{-1}$, $1 \times 10^{-4} \text{ sec}^{-1}$, $5 \times 10^{-5} \text{ sec}^{-1}$, $1 \times 10^{-5} \text{ sec}^{-1}$, $5 \times 10^{-6} \text{ sec}^{-1}$, $1 \times 10^{-6} \text{ sec}^{-1}$, $5 \times 10^{-7} \text{ sec}^{-1}$, or $1 \times 10^{-7} \text{ sec}^{-1}$.
25

In other embodiments, antibodies of the invention bind to protein tyrosine kinase biomarker polypeptides or fragments or variants thereof with an on rate (kon) of greater than or equal to $1 \times 10^3 \text{ M}^{-1} \text{ sec}^{-1}$, $5 \times 10^3 \text{ M}^{-1} \text{ sec}^{-1}$, $1 \times 10^4 \text{ M}^{-1} \text{ sec}^{-1}$, or $5 \times 10^4 \text{ M}^{-1} \text{ sec}^{-1}$. More preferably, antibodies of the invention bind to protein tyrosine
30 kinase biomarker polypeptides or fragments or variants thereof with an on rate greater than or equal to $1 \times 10^5 \text{ M}^{-1} \text{ sec}^{-1}$, $5 \times 10^5 \text{ M}^{-1} \text{ sec}^{-1}$, $1 \times 10^6 \text{ M}^{-1} \text{ sec}^{-1}$, $5 \times 10^6 \text{ M}^{-1} \text{ sec}^{-1}$, or $1 \times 10^7 \text{ M}^{-1} \text{ sec}^{-1}$.

The present invention also provides antibodies that competitively inhibit the binding of an antibody to an epitope of the invention as determined by any method known in the art for determining competitive binding, for example, the immunoassays as described herein. In preferred embodiments, the antibody competitively inhibits
5 binding to an epitope by at least 95%, at least 90%, at least 85 %, at least 80%, at least 75%, at least 70%, at least 60%, or at least 50%.

Antibodies of the present invention can act as agonists or antagonists of the protein tyrosine kinase biomarker polypeptides of the present invention. For example, the present invention includes antibodies which disrupt receptor/ligand interactions
10 with polypeptides of the invention either partially or fully. The invention includes both receptor-specific antibodies and ligand-specific antibodies. The invention also includes receptor-specific antibodies which do not prevent ligand binding, but do prevent receptor activation. Receptor activation (i.e., signaling) can be determined by techniques described herein or as otherwise known in the art. For example, receptor
15 activation can be determined by detecting the phosphorylation (e.g., on tyrosine or serine/threonine) of the receptor or its substrate by immunoprecipitation followed by Western Blot analysis. In specific embodiments, antibodies are provided that inhibit ligand activity or receptor activity by at least 95%, at least 90%, at least 85%, at least 80%, at least 75%, at least 70%, at least 60%, or at least 50% of the activity in the
20 absence of the antibody.

In another embodiment of the present invention, antibodies that immunospecifically bind to a protein tyrosine kinase biomarker, or a fragment or variant thereof, comprise a polypeptide having the amino acid sequence of any one of the heavy chains expressed by an anti-protein tyrosine kinase biomarker antibody-
25 expressing cell line of the invention, and/or any one of the light chains expressed by an anti-protein tyrosine kinase biomarker antibody-expressing cell line of the invention.

In another embodiment of the present invention, antibodies that immunospecifically bind to a tyrosine kinase biomarker protein or a fragment or variant thereof, comprise a polypeptide having the amino acid sequence of any one of
30 the V_H domains of a heavy chain expressed by an anti-protein tyrosine kinase biomarker antibody-expressing cell line, and/or any one of the V_L domains of a light

chain expressed by an anti-protein tyrosine kinase biomarker antibody-expressing cell line. In preferred embodiments, antibodies of the present invention comprise the amino acid sequence of a V_H domain and V_L domain expressed by a single anti-protein tyrosine kinase biomarker protein antibody-expressing cell line. In alternative
5 embodiments, antibodies of the present invention comprise the amino acid sequence of a V_H domain and a V_L domain expressed by two different anti-protein tyrosine kinase biomarker antibody-expressing cell lines.

Molecules comprising, or alternatively consisting of, antibody fragments or variants of the V_H and/or V_L domains expressed by an anti-protein tyrosine kinase
10 biomarker antibody-expressing cell line that immunospecifically bind to a tyrosine kinase biomarker protein, e.g., Src tyrosine kinase, are also encompassed by the invention, as are nucleic acid molecules encoding these V_H and V_L domains, molecules, fragments and/or variants.

The present invention also provides antibodies that immunospecifically bind
15 to a polypeptide, or polypeptide fragment or variant of a tyrosine kinase biomarker protein, e.g., a Src kinase biomarker protein, wherein such antibodies comprise, or alternatively consist of, a polypeptide having an amino acid sequence of any one, two, three, or more of the V_H CDRs contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody expressing cell lines. In particular,
20 the invention provides antibodies that immunospecifically bind to a tyrosine kinase biomarker protein, comprising, or alternatively consisting of, a polypeptide having the amino acid sequence of a V_H CDR1 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody expressing cell lines. In another embodiment, antibodies that immunospecifically bind to a tyrosine kinase biomarker
25 protein, comprise, or alternatively consist of, a polypeptide having the amino acid sequence of a V_H CDR2 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody expressing cell lines. In a preferred embodiment, antibodies that immunospecifically bind to a tyrosine kinase biomarker protein, comprise, or alternatively consist of, a polypeptide having the amino acid
30 sequence of a V_H CDR3 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody expressing cell line of the invention. Molecules comprising, or alternatively consisting of, these antibodies or antibody

fragments or variants thereof that immunospecifically bind to a tyrosine kinase biomarker protein or a tyrosine kinase biomarker protein fragment or variant thereof are also encompassed by the invention, as are nucleic acid molecules encoding these antibodies, molecules, fragments and/or variants.

5 The present invention also provides antibodies that immunospecifically bind to a polypeptide, or polypeptide fragment or variant of a tyrosine kinase biomarker protein, e.g., a Src kinase biomarker protein, wherein the antibodies comprise, or alternatively consist of, a polypeptide having an amino acid sequence of any one, two, three, or more of the V_L CDRs contained in a heavy chain expressed by one or more
10 anti-tyrosine kinase biomarker protein antibody expressing cell lines of the invention. In particular, the invention provides antibodies that immunospecifically bind to a tyrosine kinase biomarker protein, comprising, or alternatively consisting of, a polypeptide having the amino acid sequence of a V_L CDR1 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing
15 cell lines of the invention. In another embodiment, antibodies that immunospecifically bind to a src biomarker protein, comprise, or alternatively consist of, a polypeptide having the amino acid sequence of a V_L CDR2 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing cell lines of the invention. In a preferred embodiment, antibodies that
20 immunospecifically bind to a tyrosine kinase biomarker protein, comprise, or alternatively consist of, a polypeptide having the amino acid sequence of a V_L CDR3 contained in a heavy chain expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing cell lines of the invention. Molecules comprising, or alternatively consisting of, these antibodies, or antibody fragments or variants thereof,
25 that immunospecifically bind to a tyrosine kinase biomarker protein or a tyrosine kinase biomarker protein fragment or variant thereof are also encompassed by the invention, as are nucleic acid molecules encoding these antibodies, molecules, fragments and/or variants.

 The present invention also provides antibodies (including molecules
30 comprising, or alternatively consisting of, antibody fragments or variants) that immunospecifically bind to a tyrosine kinase biomarker protein, polypeptide or polypeptide fragment or variant of a tyrosine kinase biomarker protein, e.g., Src

tyrosine kinase, wherein the antibodies comprise, or alternatively consist of, one, two, three, or more V_H CDRs, and one, two, three or more V_L CDRs, as contained in a heavy chain or light chain expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing cell lines of the invention. In particular, the invention provides antibodies that immunospecifically bind to a polypeptide or polypeptide fragment or variant of a tyrosine kinase biomarker protein, wherein the antibodies comprise, or alternatively consist of, a V_H CDR1 and a V_L CDR1, a V_H CDR1 and a V_L CDR2, a V_H CDR1 and a V_L CDR3, a V_H CDR2 and a V_L CDR1, a V_H CDR2 and a V_L CDR2, a V_H CDR2 and a V_L CDR3, a V_H CDR3 and a V_H CDR1, a V_H CDR3 and a V_L CDR2, a V_H CDR3 and a V_L CDR3, or any combination thereof, of the V_H CDRs and V_L CDRs contained in a heavy chain or light chain immunoglobulin molecule expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing cell lines of the invention. In a preferred embodiment, one or more of these combinations are from a single anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention. Molecules comprising, or alternatively consisting of, fragments or variants of these antibodies that immunospecifically bind to the tyrosine kinase biomarker proteins are also encompassed by the invention, as are nucleic acid molecules encoding these antibodies, molecules, fragments or variants.

The present invention also provides nucleic acid molecules, generally isolated, encoding an antibody of the invention (including molecules comprising, or alternatively consisting of, antibody fragments or variants thereof). In a specific embodiment, a nucleic acid molecule of the invention encodes an antibody (including molecules comprising, or alternatively consisting of, antibody fragments or variants thereof), comprising, or alternatively consisting of, a V_H domain having an amino acid sequence of any one of the V_H domains of a heavy chain expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention and a V_L domain having an amino acid sequence of a light chain expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention. In another embodiment, a nucleic acid molecule of the invention encodes an antibody (including molecules comprising, or alternatively consisting of, antibody fragments or variants thereof), comprising, or alternatively consisting of, a V_H domain having an

amino acid sequence of any one of the V_H domains of a heavy chain expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention, or a V_L domain having an amino acid sequence of a light chain expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention.

5 The present invention also provides antibodies that comprise, or alternatively consist of, variants (including derivatives) of the antibody molecules (e.g., the V_H domains and/or V_L domains) described herein, which antibodies immunospecifically bind to a tyrosine kinase biomarker protein or fragment or variant thereof, e.g., a Src tyrosine kinase polypeptide.

10 Standard techniques known to those of skill in the art can be used to introduce mutations in the nucleotide sequence encoding a molecule of the invention, including, for example, site-directed mutapolynucleotidesis and PCR-mediated mutapolynucleotidesis which result in amino acid substitutions. Preferably the molecules are immunoglobulin molecules. Also preferably, the variants (including
15 derivatives) encode less than 50 amino acid substitutions, less than 40 amino acid substitutions, less than 30 amino acid substitutions, less than 25 amino acid substitutions, less than 20 amino acid substitutions, less than 15 amino acid substitutions, less than 10 amino acid substitutions, less than 5 amino acid substitutions, less than 4 amino acid substitutions, less than 3 amino acid
20 substitutions, or less than 2 amino acid substitutions, relative to the reference V_H domain, V_H CDR1, V_H CDR2, V_H CDR3, V_L domain, V_L CDR1, V_L CDR2, or V_L CDR3.

 A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a side chain with a similar
25 charge. Families of amino acid residues having side chains with similar charges have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine,
30 proline, phenylalanine, methionine, tryptophan), beta-branched side chains (e.g., threonine, valine, isoleucine) and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine). Alternatively, mutations can be introduced randomly along all

or part of the coding sequence, such as by saturation mutapolynucleotidesis. The resultant mutants can be screened for biological activity to identify mutants that retain activity.

For example, it is possible to introduce mutations only in framework regions or only in CDR regions of an antibody molecule. Introduced mutations can be silent or neutral missense mutations, i.e., have no, or little, effect on an antibody's ability to bind antigen. These types of mutations can be useful to optimize codon usage, or to improve hybridoma antibody production. Alternatively, non-neutral missense mutations can alter an antibody's ability to bind antigen. The location of most silent and neutral missense mutations is likely to be in the framework regions, while the location of most non-neutral missense mutations is likely to be in the CDRs, although this is not an absolute requirement. One of skill in the art is able to design and test mutant molecules with desired properties, such as no alteration in antigen binding activity or alteration in binding activity (e.g., improvements in antigen binding activity or change in antibody specificity). Following mutapolynucleotidesis, the encoded protein may routinely be expressed and the functional and/or biological activity of the encoded protein can be determined using techniques described herein or by routinely modifying techniques known and practiced in the art.

In a specific embodiment, an antibody of the invention (including a molecule comprising, or alternatively consisting of, an antibody fragment or variant thereof), that immunospecifically binds to protein tyrosine kinase biomarker polypeptides or fragments or variants thereof, comprises, or alternatively consists of, an amino acid sequence encoded by a nucleotide sequence that hybridizes to a nucleotide sequence that is complementary to that encoding one of the V_H or V_L domains expressed by one or more anti-tyrosine kinase biomarker protein antibody-expressing cell lines of the invention, preferably under stringent conditions, e.g., hybridization to filter-bound DNA in 6x sodium chloride/sodium citrate (SSC) at about 45°C followed by one or more washes in 0.2 x SSC/0.1% SDS at about 50°C-65°C, preferably under highly stringent conditions, e.g., hybridization to filter-bound nucleic acid in 6xSSC at about 45°C followed by one or more washes in 0.1xSSC/0.2% SDS at about 68°C, or under other stringent hybridization conditions which are known to those of skill in the art

(see, for example, Ausubel, F.M. et al., eds., 1989, Current Protocols in Molecular Biology, Vol. I, Green Publishing Associates, Inc. and John Wiley & Sons, Inc., New York at pages 6.3.1-6.3.6 and 2.10.3). Nucleic acid molecules encoding these antibodies are also encompassed by the invention.

5 It is well known within the art that polypeptides, or fragments or variants thereof, with similar amino acid sequences often have similar structure and many of the same biological activities. Thus, in one embodiment, an antibody (including a molecule comprising, or alternatively consisting of, an antibody fragment or variant thereof), that immunospecifically binds to a protein tyrosine kinase biomarker
10 polypeptide or fragments or variants of a tyrosine kinase biomarker polypeptide, comprises, or alternatively consists of, a V_H domain having an amino acid sequence that is at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 99% identical to the amino acid sequence of a V_H domain of a
15 heavy chain expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention.

 In another embodiment, an antibody (including a molecule comprising, or alternatively consisting of, an antibody fragment or variant thereof), that immunospecifically binds to a tyrosine kinase biomarker polypeptide, or fragments or
20 variants of a tyrosine kinase biomarker protein polypeptide, comprises, or alternatively consists of, a V_L domain having an amino acid sequence that is at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 99% identical to the amino acid sequence of a V_L domain of a light chain
25 expressed by an anti-tyrosine kinase biomarker protein antibody-expressing cell line of the invention.

 The present invention also provides antibodies (including molecules comprising, or alternatively consisting of, antibody fragments or variants thereof), that down-regulate the cell-surface expression of a tyrosine kinase biomarker protein, as
30 determined by any method known in the art such as, for example, FACS analysis or immunofluorescence assays. By way of a non-limiting hypothesis, such down-regulation can be the result of antibody-induced internalization of a tyrosine kinase

biomarker protein. Such antibodies can comprise, or alternatively consist of, a portion (e. g., V_H CDR1, V_H CDR2, V_H CDR3, V_L CDR1, V_L CDR2, or V_L CDR3) of a V_H or V_L domain having an amino acid sequence of an antibody of the invention, or a fragment or variant thereof.

5 In another embodiment, an antibody that down-regulates the cell-surface expression of a tyrosine kinase biomarker protein comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_H domain of an antibody of the invention, or a fragment or variant thereof and a V_L domain of an antibody of the invention, or a fragment or variant thereof. In another embodiment, an antibody that
10 down-regulates the cell-surface expression of a tyrosine kinase biomarker protein comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_H domain and a V_L domain from a single antibody (or scFv or Fab fragment) of the invention, or fragments or variants thereof. In another embodiment, an antibody that down-regulates the cell-surface expression of a tyrosine kinase biomarker protein
15 comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_H domain of an antibody of the invention, or a fragment or variant thereof. In another embodiment, an antibody that down-regulates the cell-surface expression of a tyrosine kinase biomarker protein comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_L domain of an antibody of the
20 invention, or a fragment or variant thereof.

In a preferred embodiment, an antibody that down-regulates the cell-surface expression of a tyrosine kinase biomarker protein comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_H CDR3 of an antibody of the invention, or a fragment or variant thereof. In another preferred embodiment, an
25 antibody that down-regulates the cell-surface expression of a tyrosine kinase biomarker protein comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_L CDR3 of an antibody of the invention, or a fragment or variant thereof. Nucleic acid molecules encoding these antibodies are also encompassed by the invention.

30 In another preferred embodiment, an antibody that enhances the activity of a tyrosine kinase biomarker protein, or a fragment or variant thereof, comprises, or alternatively consists of, a polypeptide having the amino acid sequence of a V_L CDR3

of an antibody of the invention, or a fragment or variant thereof. Nucleic acid molecules encoding these antibodies are also encompassed by the invention.

As nonlimiting examples, antibodies of the present invention can be used to purify, detect, and target the protein tyrosine kinase polypeptides of the present invention, including both *in vitro* and *in vivo* diagnostic, detection, screening, and/or therapeutic methods. For example, the antibodies have been used in immunoassays for qualitatively and quantitatively measuring levels of src biomarker polypeptides in biological samples. (See, e.g., Harlow et al., *Antibodies : A Laboratory Manual*, Cold Spring Harbor Laboratory Press, 2nd Ed. 1988, which is incorporated by reference herein in its entirety).

By way of another nonlimiting example, antibodies of the invention can be administered to individuals as a form of passive immunization. Alternatively, antibodies of the present invention can be used for epitope mapping to identify the epitope(s) that are bound by the antibody. Epitopes identified in this way can, in turn, for example, be used as vaccine candidates, i.e., to immunize an individual to elicit antibodies against the naturally-occurring forms of one or more tyrosine kinase biomarker proteins.

As discussed in more detail below, the antibodies of the present invention can be used either alone or in combination with other compositions. The antibodies can further be recombinantly fused to a heterologous polypeptide at the N-or C-terminus, or chemically conjugated (including covalent and non-covalent conjugations) to polypeptides or other compositions. For example, antibodies of the present invention can be recombinantly fused or conjugated to molecules that are useful as labels in detection assays and to effector molecules such as heterologous polypeptides, drugs, radionuclides, or toxins. See, e.g., PCT publications WO 92/08495; WO 91/14438; WO 89/12624; U.S. Patent No. 5,314,995 and EP 396, 387.

The antibodies of the invention include derivatives that are modified, i.e., by the covalent attachment of any type of molecule to the antibody. For example, without limitation, the antibody derivatives include antibodies that have been modified, e.g., by glycosylation, acetylation, pegylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc. Any of numerous chemical modifications can be

carried out by known techniques, including, but not limited to, specific chemical cleavage, acetylation, formylation, metabolic synthesis of tunicamycin, and the like. In addition, the antibody derivative can contain one or more non-classical amino acids.

5 The antibodies of the present invention can be generated by any suitable method known in the art. Polyclonal antibodies directed against an antigen or immunogen of interest can be produced by various procedures well known in the art. For example, a tyrosine kinase biomarker polypeptide or peptide of the invention can be administered to various host animals as elucidated above to induce the production
10 of sera containing polyclonal antibodies specific for the biomarker antigen. Various adjuvants can also be used to increase the immunological response, depending on the host species; adjuvants include, but are not limited to, Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet
15 hemocyanins, dinitrophenol, and potentially useful human adjuvants such as BCG (bacille Calmette-Guerin) and corynebacterium parvum. Such adjuvants are also well known in the art.

Monoclonal antibodies can be prepared using a wide variety of techniques known in the art, including the use of hybridoma, recombinant and phage display
20 technologies, or a combination thereof. For example, monoclonal antibodies can be produced using hybridoma techniques as known and practiced in the art and as taught, for example, in Kohler and Milstein, 1975, *Nature*, 256:495-497; Harlow et al., *Antibodies : A Laboratory Manual*, Cold Spring Harbor Laboratory Press, 2nd Ed. 1988; and Hammerling, et al., In: *Monoclonal Antibodies and T-Cell Hybridomas*,
25 Elsevier, N. Y., pages 563-681, 1981, the contents of which are incorporated herein by reference in their entireties. The term "monoclonal antibody" as used herein is not limited to antibodies produced through hybridoma technology. The term "monoclonal antibody" refers to an antibody that is derived from a single clone, including any eukaryotic, prokaryotic, or phage clone, and does not necessarily refer to the method
30 by which it is produced. Techniques involving continuous cell line cultures can also be used. In addition to the hybridoma technique, other techniques include the trioma technique, the human B-cell hybridoma technique (Kozbor et al., 1983, *Immunol.*

Today, 4:72), and the EBV-hybridoma technique (Cole et al., 1985. In: *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc., pp. 77-96).

Methods for producing and screening for specific antibodies using hybridoma technology are routine and well known in the art. As a nonlimiting example, mice can be immunized with a tyrosine kinase polypeptide or peptide of the invention, or variant thereof, or with a cell expressing the polypeptide or peptide or variant thereof. Once an immune response is detected, e.g., antibodies specific for the antigen are detected in the sera of immunized mice, the spleen is harvested and splenocytes are isolated. The splenocytes are then fused by well known techniques to any suitable myeloma cells, for example cells from cell line SP2/0 or P3X63-AG8.653 available from the ATCC. Hybridomas are selected and cloned by limiting dilution techniques. The hybridoma clones are then assayed by methods known in the art to determine and select those cells that secrete antibodies capable of binding to a polypeptide of the invention. Ascites fluid, which generally contains high levels of antibodies, can be generated by immunizing mice with positive hybridoma clones.

Another well known method for producing both polyclonal and monoclonal human B cell lines is transformation using Epstein Barr Virus (EBV). Protocols for generating EBV-transformed B cell lines are commonly known in the art, such as, for example, the protocol outlined in Chapter 7.22 of *Current Protocols in Immunology*, Coligan et al., Eds., 1994, John Wiley & Sons, NY, which is hereby incorporated by reference herein in its entirety. The source of B cells for transformation is commonly human peripheral blood, but B cells for transformation can also be obtained from other sources including, but not limited to, lymph node, tonsil, spleen, tumor tissue, and infected tissues. Tissues are generally prepared as single cell suspensions prior to EBV transformation. In addition, T cells that may be present in the B cell samples can be either physically removed or inactivated (e.g., by treatment with cyclosporin A). The removal of T cells is often advantageous, because T cells from individuals seropositive for anti-EBV antibodies can suppress B cell immortalization by EBV. In general, a sample containing human B cells is innoculated with EBV and cultured for 3-4 weeks. A typical source of EBV is the culture supernatant of the B95-8 cell line (ATCC; VR-1492). Physical signs of EBV transformation can generally be seen toward the end of the 3-4 week culture period.

By phase-contrast microscopy, transformed cells appear large, clear and "hairy"; they tend to aggregate in tight clusters of cells. Initially, EBV lines are generally polyclonal. However, over prolonged periods of cell culture, EBV lines can become monoclonal as a result of the selective outgrowth of particular B cell clones.

5 Alternatively, polyclonal EBV transformed lines can be subcloned (e.g., by limiting dilution) or fused with a suitable fusion partner and plated at limiting dilution to obtain monoclonal B cell lines. Suitable fusion partners for EBV transformed cell lines include mouse myeloma cell lines (e.g., SP2/0, X63-Ag8.653), heteromyeloma cell lines (human x mouse ; e.g., SPAM-8, SBC-H20, and CB-F7), and human cell

10 lines (e.g., GM 1500, SKO-007, RPMI 8226, and KR-4). Thus, the present invention also includes a method of generating polyclonal or monoclonal human antibodies against protein tyrosine kinase polypeptides and peptides of the invention, or fragments thereof, comprising EBV-transformation of human B cells.

Antibody fragments that recognize specific epitopes can be generated by

15 known techniques. For example, Fab and F(ab')₂ fragments of the invention can be produced by proteolytic cleavage of immunoglobulin molecules, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F (ab')₂ fragments). F(ab')₂ fragments contain the variable region, the light chain constant region and the CH1 domain of the heavy chain.

20 Antibodies encompassed by the present invention can also be generated using various phage display methods known in the art. In phage display methods, functional antibody domains are displayed on the surface of phage particles which carry the polynucleotide sequences encoding them. In a particular embodiment, such phage can be utilized to display antigen binding domains expressed from a repertoire or

25 combinatorial antibody library (e.g., human or murine). Phage expressing an antigen binding domain that binds to the antigen of interest can be selected or identified with antigen, e.g., using labeled antigen or antigen bound or captured onto a solid surface or bead. Phage used in these methods are typically filamentous phage including fd and M13 binding domains expressed from phage with Fab, Fv, or disulfide stabilized

30 antibody domains recombinantly fused to either the phage polynucleotide III or polynucleotide VIII protein. Examples of phage display methods that can be used to make the antibodies of the present invention include those disclosed in Brinkman et

al., 1995, *J. Immunol. Methods*, 182:41-50; Ames et al., 1995, *J. Immunol. Methods*, 184:177-186; Kettleborough et al., 1994, *Eur. J. Immunol.*, 24:952-958; Persic et al., 1997, *Gene*, 187:9-18; Burton et al., 1994, *Advances in Immunology*, 57:191-280; PCT application No. PCT/GB91/01134; PCT publications WO 90/02809; WO
5 91/10737; WO 92/01047; WO 92/18619; WO 93/11236; WO 95/15982; WO 95/20401; and U.S. Patent Nos. 5,698,426; 5,223,409; 5,403,484; 5,580,717; 5,427,908; 5,750,753; 5,821,047; 5,571,698; 5,427,908; 5,516,637; 5,780,225; 5,658,727; 5,733,743 and 5,969,108, each of which is incorporated herein by reference in its entirety.

10 As described in the above references, after phage selection, the antibody coding regions from the phage can be isolated and used to generate whole antibodies, including human antibodies, or any other desired antigen binding fragment, and expressed in any desired host, including mammalian cells, insect cells, plant cells, yeast, and bacteria, e.g., as described in detail below.

15 Examples of techniques that can be used to produce single-chain Fvs and antibodies include those described in U.S. Patent Nos. 4,946,778 and 5,258,498; Huston et al., 1991, *Methods in Enzymology*, 203:46-88; Shu et al., 1993, *Proc. Natl. Acad. Sci. USA*, 90:7995-7999; and Skerra et al., 1988, *Science*, 240:1038-1040. For some uses, including the *in vivo* use of antibodies in humans and in *in vitro* detection
20 assays, it may be preferable to use chimeric, humanized, or human antibodies. A chimeric antibody is a molecule in which different portions of the antibody are derived from different animal species, such as antibodies having a variable region derived from a murine monoclonal immunoglobulin and a human immunoglobulin constant region. Methods for producing chimeric antibodies are known in the art.
25 (See, e.g., Morrison, 1985, *Science*, 229:1202; Oi et al., 1986, *BioTechniques*, 4:214; Gillies et al., 1989, *J. Immunol. Methods*, 125:191-202; and U.S. Patent Nos. 5,807,715; 4,816,567; and 4,816,397, which are incorporated herein by reference in their entirety).

Humanized antibodies are antibody molecules from non-human species that
30 bind to the desired antigen and have one or more complementarity determining regions (CDRs) from the nonhuman species and framework regions from a human immunoglobulin molecule. Often, framework residues in the human framework

regions are substituted with corresponding residues from the CDR and framework regions of the donor antibody to alter, preferably improve, antigen binding. These framework substitutions are identified by methods well known in the art, e.g., by modeling of the interactions of the CDR and framework residues to identify framework residues important for antigen binding, and by sequence comparison to identify unusual framework residues at particular positions. (See, e.g., Queen et al., U.S. Patent Nos. 5,693,762 and 5,585,089; and Riechmann et al., 1988, *Nature*, 332:323, which are incorporated herein by reference in their entireties). Antibodies can be humanized using a variety of techniques known in the art, including, for example, CDR-grafting (EP 239,400; PCT publication WO 91/09967; U.S. Patent Nos. 5,225,539; 5,530,101; and 5,585,089); veneering or resurfacing (EP 592,106; EP 519,596; Padlan, 1991, *Molecular Immunology*, 28(4/5):489-498; Studnicka et al., 1994, *Protein Engineering*, 7(6):805-814; Roguska et al., 1994, *Proc. Natl. Acad. Sci. USA*, 91:969-973; and chain shuffling (U.S. Patent No. 5,565,332).

Completely human antibodies can be made by a variety of methods known in the art, including the phage display methods described above, using antibody libraries derived from human immunoglobulin sequences. See also, U.S. Patent Nos. 4,444,887 and 4,716,111; and PCT publications WO 98/46645, WO 98/50433, WO 98/24893, WO 98/16654, WO 96/34096, WO 96/33735, and WO 91/10741; each of which is incorporated herein by reference in its entirety.

Completely human antibodies are particularly desirable for therapeutic treatment of human patients, so as to avoid or alleviate immune reaction to foreign protein. Human antibodies can be made by a variety of methods known in the art, including the phage display methods described above, using antibody libraries derived from human immunoglobulin sequences. See also, U.S. Patent Nos. 4,444,887 and 4,716,111; and PCT publications WO 98/46645, WO 98/50433, WO 98/24893, WO 98/16654, WO 96/34096, WO 96/33735, and WO 91/10741; each of which is incorporated herein by reference in its entirety.

Human antibodies can also be produced using transgenic mice which are incapable of expressing functional endogenous immunoglobulins, but which can express human immunoglobulin polynucleotides. For example, the human heavy and light chain immunoglobulin polynucleotide complexes can be introduced randomly,

or by homologous recombination, into mouse embryonic stem cells. Alternatively, the human variable region, constant region, and diversity region may be introduced into mouse embryonic stem cells, in addition to the human heavy and light chain polynucleotides. The mouse heavy and light chain immunoglobulin polynucleotides
5 can be rendered nonfunctional separately or simultaneously with the introduction of human immunoglobulin loci by homologous recombination. In particular, homozygous deletion of the J_H region prevents endogenous antibody production. The modified embryonic stem cells are expanded and microinjected into blastocysts to produce chimeric mice. The chimeric mice are then bred to produce homozygous
10 offspring which express human antibodies. The transgenic mice are immunized in the normal fashion with a selected antigen, e.g., all or a portion of a polypeptide of the invention.

Thus, using such a technique, it is possible to produce useful human IgG, IgA, IgM, IgD and IgE antibodies. For an overview of the technology for producing
15 human antibodies, see Lonberg and Huszar, 1995, *Intl. Rev. Immunol.*, 13:65-93. For a detailed discussion of the technology for producing human antibodies and human monoclonal antibodies and protocols for producing such antibodies, see, e.g., PCT publications WO 98/24893; WO 92/01047; WO 96/34096; WO 96/33735; European Patent No. 0 598 877; U.S. Patent Nos. 5,413,923; 5,625,126; 5,633,425; 5,569,825;
20 5,661,016; 5,545,806; 5,814,318; 5,885,793; 5,916,771; 5,939,598; 6,075,181; and 6,114,598, which are incorporated by reference herein in their entirety. In addition, companies such as Abgenix, Inc. (Fremont, CA), Protein Design Labs, Inc. (Mountain View, CA) and Genpharm (San Jose, CA) can be engaged to provide human antibodies directed against a selected antigen using technology similar to the above
25 described technologies.

Completely human antibodies which recognize a selected epitope can be generated using a technique referred to as "guided selection". In this approach, a selected non-human monoclonal antibody, e.g., a mouse antibody, is used to guide the selection of a completely human antibody recognizing the same epitope. (Jespers et
30 al., 1988, *BioTechnology*, 12:899-903).

Further, antibodies to the protein tyrosine kinase polypeptides of the invention can, in turn, be utilized to generate anti-idiotypic antibodies that "mimic" protein

tyrosine kinase biomarker polypeptides of the invention using techniques well known to those skilled in the art. (See, e.g., Greenspan and Bona, 1989, *FASEB J.*, 7(5):437-444 and Nissinoff, 1991, *J. Immunol.*, 147(8):2429-2438). For example, antibodies which bind to and competitively inhibit polypeptide multimerization and/or binding of
5 a polypeptide of the invention to a ligand can be used to generate anti-idiotypes that "mimic" the polypeptide multimerization and/or binding domain and, as a consequence, bind to and neutralize the polypeptide and/or its ligand, e.g., in therapeutic regimens. Such neutralizing anti-idiotypes or Fab fragments of such anti-idiotypes can be used to neutralize polypeptide ligand. For example, such anti-
10 idiotypic antibodies can be used to bind a polypeptide of the invention and/or to bind its ligands/receptors, and thereby activate or block its biological activity.

Intrabodies are antibodies, often scFvs, that are expressed from a recombinant nucleic acid molecule and are engineered to be retained intracellularly (e.g., retained in the cytoplasm, endoplasmic reticulum, or periplasm of the host cells). Intrabodies
15 can be used, for example, to ablate the function of a protein to which the intrabody binds. The expression of intrabodies can also be regulated through the use of inducible promoters in the nucleic acid expression vector comprising nucleic acid encoding the intrabody. Intrabodies of the invention can be produced using methods known in the art, such as those disclosed and reviewed in Chen et al., 1994, *Hum.*
20 *Polynucleotide Ther.*, 5:595-601; Marasco, W.A., 1997, *Polynucleotide Ther.*, 4:11-15; Rondon and Marasco, 1997, *Annu. Rev. Microbiol.*, 51:257-283; Proba et al., 1998, *J. Mol. Biol.*, 275:245-253; Cohen et al., 1998, *Oncogene*, 17:2445-2456; Ohage and Steipe, 1999, *J. Mol. Biol.*, 291:1119-1128; Ohage et al., 1999, *J. Mol. Biol.*, 291:1129-1134; Wirtz and Steipe, 1999, *Protein Sci.*, 8:2245-2250; and Zhu et
25 al., 1999, *J. Immunol. Methods*, 231:207-222.

XenoMouse Technology Antibodies in accordance with the invention are preferably prepared by the utilization of a transgenic mouse that has a substantial portion of the human antibody producing genome inserted into its genome, but that is rendered deficient in the production of endogenous murine antibodies (e.g.,
30 XenoMouse strains available from Abgenix Inc., Fremont, CA). Such mice are capable of producing human immunoglobulin molecules and are virtually deficient in the production of murine immunoglobulin molecules. Technologies utilized for

achieving the same are disclosed in the patents, applications, and references disclosed herein.

The ability to clone and reconstruct megabase-sized human loci in YACs and to introduce them into the mouse germline provides a powerful approach to elucidating the functional components of very large or crudely mapped loci, as well as generating useful models of human disease. Furthermore, the utilization of such technology for substitution of mouse loci with their human equivalents could provide unique insights into the expression and regulation of human polynucleotide products during development, their communication with other systems, and their involvement in disease induction and progression. An important practical application of such a strategy is the "humanization" of the mouse humoral immune system. Introduction of human immunoglobulin (Ig) loci into mice in which the endogenous Ig polynucleotides have been inactivated offers the opportunity to study the mechanisms underlying programmed expression and assembly of antibodies as well as their role in B cell development. Furthermore, such a strategy could provide an ideal source for the production of fully human monoclonal antibodies (Mabs), which is an important milestone toward fulfilling the promise of antibody therapy in human disease.

Fully human antibodies are expected to minimize the immunogenic and allergic responses intrinsic to mouse or mouse-derivatized monoclonal antibodies and thus to increase the efficacy and safety of the administered antibodies. The use of fully human antibodies can be expected to provide a substantial advantage in the treatment of chronic and recurring human diseases, such as cancer, which require repeated antibody administrations.

One approach toward this goal was to engineer mouse strains deficient in mouse antibody production to harbor large fragments of the human Ig loci in anticipation that such mice would produce a large repertoire of human antibodies in the absence of mouse antibodies. Large human Ig fragments would preserve the large variable polynucleotide diversity as well as the proper regulation of antibody production and expression. By exploiting the mouse machinery for antibody diversification and selection and the lack of immunological tolerance to human proteins, the reproduced human antibody repertoire in these mouse strains should yield high affinity antibodies against any antigen of interest, including human

antigens. Using the hybridoma technology, antigen-specific human monoclonal antibodies with the desired specificity could be readily produced and selected.

This general strategy was demonstrated in connection with the generation of the first "XenoMouseT" strains as published in 1994. See Green et al., 1994, *Nature Genetics*, 7:13-21. The XenoMouse strains were engineered with yeast artificial chromosomes (YACS) containing 245 kb and 10, 190 kb-sized germline configuration fragments of the human heavy chain locus and kappa light chain locus, respectively, which contained core variable and constant region sequences. *Id.* The human Ig-containing YACs proved to be compatible with the mouse system for both rearrangement and expression of antibodies and were capable of substituting for the inactivated mouse Ig polynucleotides. This was demonstrated by their ability to induce B-cell development, to produce an adult-like human repertoire of fully human antibodies, and to generate antigen-specific human monoclonal antibodies. These results also suggested that introduction of larger portions of the human Ig loci containing greater numbers of V polynucleotides, additional regulatory elements, and human Ig constant regions might recapitulate substantially the full repertoire that is characteristic of the human humoral response to infection and immunization. The work of Green et al. was recently extended to the introduction of greater than approximately 80% of the human antibody repertoire through the use of megabase-sized, germline configuration YAC fragments of the human heavy chain loci and kappa light chain loci, respectively, to produce XenoMouse mice. See Mendez et al., 1997, *Nature Genetics*, 15:146-156; Green and Jakobovits, 1998, *J. Exp. Med.*, 188:483-495; and Green, 1999, *Journal of Immunological Methods*, 231:11-23, the disclosures of which are hereby incorporated herein by reference.

Human anti-mouse antibody (HAMA) responses have led the industry to prepare chimeric or otherwise humanized antibodies. While chimeric antibodies typically are comprised of a human constant region and a murine variable region, it is expected that certain human anti-chimeric antibody (HACA) responses will be observed, particularly in treatments involving chronic or multi-dose utilizations of the antibody. Thus, it is desirable to provide fully human antibodies against protein tyrosine kinase biomarker polypeptides in order to vitiate concerns and/or effects of HAMA or HACA responses.

Antibodies of the invention can be chemically synthesized or produced through the use of recombinant expression systems. Accordingly, the invention further embraces polynucleotides comprising a nucleotide sequence encoding an antibody of the invention and fragments thereof. The invention also encompasses
5 polynucleotides that hybridize under stringent or lower stringency hybridization conditions, e.g., as defined supra, to polynucleotides that encode an antibody, preferably, an antibody that specifically binds to a protein tyrosine kinase polypeptide of this invention, and more preferably, an antibody that binds to a polypeptide having the amino acid sequence of one or more of the protein tyrosine kinase biomarker
10 sequences, e.g., Src tyrosine kinase biomarkers, as set forth in Table 2.

The polynucleotides may be obtained, and the nucleotide sequence of the polynucleotides determined, by any method known in the art. For example, if the nucleotide sequence of the antibody is known, a polynucleotide encoding the antibody can be assembled from chemically synthesized oligonucleotides (e.g., as described in
15 Kutmeier et al., 1994, *BioTechniques*, 17:242), which, briefly, involves the synthesis of overlapping oligonucleotides containing portions of the sequence encoding the antibody, the annealing and ligating of those oligonucleotides, and then the amplification of the ligated oligonucleotides by PCR.

Alternatively, a polynucleotide encoding an antibody can be generated from
20 nucleic acid from a suitable source. If a clone containing a nucleic acid encoding a particular antibody is not available, but the sequence of the antibody molecule is known, a nucleic acid encoding the immunoglobulin can be chemically synthesized or obtained from a suitable source (e.g., an antibody cDNA library, or a cDNA library generated from, (or a nucleic acid, preferably poly A+ RNA, isolated from), any tissue
25 or cells expressing the antibody, such as hybridoma cells selected to express an antibody of the invention by PCR amplification using synthetic primers hybridizable to the 3' and 5' ends of the sequence. Alternatively, cloning using an oligonucleotide probe specific for the particular polynucleotide sequence to be identified, e.g., a cDNA clone from a cDNA library that encodes the desired antibody can be employed.
30 Amplified nucleic acids generated by PCR can then be cloned into replicable cloning vectors using any method well known in the art.

Once the nucleotide sequence and corresponding encoded amino acid sequence of the antibody are determined, the nucleotide sequence of the antibody can be manipulated using methods well known in the art for the manipulation of nucleotide sequences, e.g., recombinant DNA techniques, site directed mutapolynucleotidesis, PCR, etc. (see, for example, the techniques described in
5 Sambrook et al., 1990, *Molecular Cloning, A Laboratory Manual*, 2nd Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY; and F.M. Ausubel et al., eds., 1998, *Current Protocols in Molecular Biology*, John Wiley & Sons, NY, which are both incorporated by reference herein in their entireties), to generate antibodies having
10 a different amino acid sequence, for example, to create amino acid substitutions, deletions, and/or insertions.

In a specific embodiment, the amino acid sequence of the heavy and/or light chain variable domains can be inspected to identify the sequences of the CDRs by methods that are well known in the art, e.g., by comparison to known amino acid
15 sequences of other heavy and light chain variable regions, to determine the regions of sequence hypervariability. Using routine recombinant DNA techniques, one or more of the CDRs can be inserted within framework regions, e.g., into human framework regions, to humanize a non-human antibody, as described *supra*. The framework regions can be naturally occurring or consensus framework regions, and preferably,
20 are human framework regions (see, e.g., Chothia et al., 1998, *J. Mol. Biol.*, 278:457-479, for a listing of human framework regions).

Preferably, the polynucleotide generated by the combination of the framework regions and CDRs encodes an antibody that specifically binds to a protein tyrosine kinase biomarker polypeptide of the invention. Also preferably, as discussed *supra*,
25 one or more amino acid substitutions can be made within the framework regions; such amino acid substitutions are performed with the goal of improving binding of the antibody to its antigen, e.g., greater antibody binding affinity. In addition, such methods can be used to make amino acid substitutions or deletions of one or more variable region cysteine residues participating in an intrachain disulfide bond to
30 generate antibody molecules lacking one or more intrachain disulfide bonds. Other alterations that can be made to the polynucleotide are encompassed by the present invention and are within the skill of the art.

For some uses, such as for *in vitro* affinity maturation of an antibody of the invention, it is useful to express the V_H and V_L domains of the heavy and light chains of one or more antibodies of the invention as single chain antibodies, or Fab fragments, in a phage display library using phage display methods as described *supra*.

5 For example, the cDNAs encoding the V_H and V_L domains of one or more antibodies of the invention can be expressed in all possible combinations using a phage display library, thereby allowing for the selection of V_H/V_L combinations that bind to the protein tyrosine kinase biomarker polypeptides according to the present invention with preferred binding characteristics such as improved affinity or improved off rates.

10 In addition, V_H and V_L segments, particularly, the CDR regions of the V_H and V_L domains of one or more antibodies of the invention, can be mutated *in vitro*. Expression of V_H and V_L domains with "mutant" CDRs in a phage display library allows for the selection of V_H/V_L combinations that bind to protein tyrosine kinase biomarkers, e.g., Src tyrosine kinase biomarker proteins, which are receptor

15 polypeptides with preferred binding characteristics such as improved affinity or improved off rates.

Once an antibody molecule of the invention has been produced by an animal, chemically synthesized, or recombinantly expressed, it can be purified by any method known in the art for the purification of an immunoglobulin or polypeptide molecule,

20 for example, by chromatography (e.g., ion exchange, affinity, particularly by affinity for the specific antigen, Protein A, and sizing column chromatography), centrifugation, differential solubility, or by any other standard technique for the purification of proteins. In addition, the antibodies of the present invention or fragments thereof can be fused to heterologous polypeptide sequences described

25 herein or otherwise known in the art, to facilitate purification.

The present invention encompasses antibodies that are recombinantly fused or chemically conjugated (including both covalently and non-covalently conjugated) to a polypeptide (or portion thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino acids of the polypeptide) of the present invention to generate fusion

30 proteins. The fusion does not necessarily need to be direct, but can occur through linker sequences. The antibodies can be specific for antigens other than polypeptides (or portions thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino

acids of the polypeptide) of the present invention. For example, antibodies can be used to target the polypeptides of the present invention to particular cell types, either *in vitro* or *in vivo*, by fusing or conjugating the polypeptides of the present invention to antibodies specific for particular cell surface receptors.

5 The present invention further includes compositions comprising the protein tyrosine kinase biomarker polypeptides of the present invention fused or conjugated to antibody domains other than the variable region domain. For example, the polypeptides of the present invention can be fused or conjugated to an antibody Fc region, or portion thereof. The antibody portion fused to a polypeptide of the present
10 invention can comprise the constant region, hinge region, CH1 domain, CH2 domain, CH3 domain, or any combination of whole domains or portions thereof. The polypeptides can also be fused or conjugated to the above antibody portions to form multimers. For example, Fc portions of immunoglobulin molecules fused to the polypeptides of the present invention can form dimers through disulfide bonding
15 between the Fc portions. Higher multimeric forms can be made by fusing the polypeptides to portions of IgA and IgM. Methods for fusing or conjugating the polypeptides of the present invention to antibody portions are known in the art. (See, e.g., U.S. Patent Nos. 5,336,603; 5,622,929; 5,359,046; 5,349,053; 5,447,851; 5,112,946; EP 307,434; EP 367,166; PCT publications WO 96/04388; WO 91/06570;
20 Ashkenazi et al., 1991, *Proc. Natl. Acad. Sci. USA*, 88:10535-10539; Zheng et al., 1995, *J. Immunol.*, 154:5590-5600; and Vil et al., *Proc. Natl. Acad. Sci. USA*, 89:11337-11341, which are hereby incorporated by reference herein in their entireties).

 As discussed *supra*, the polypeptides corresponding to a polypeptide,
25 polypeptide fragment, or a variant of one or more of the protein tyrosine kinase biomarker amino acid sequences as set forth in Table 2 can be fused or conjugated to the above antibody portions to increase the *in vivo* half life of the polypeptides, or for use in immunoassays using methods known in the art. Further, the polypeptides corresponding to one or more of the protein tyrosine kinase biomarker, e.g., src
30 biomarker, sequences as set forth in Table 2 can be fused or conjugated to the above antibody portions to facilitate purification. For guidance, chimeric proteins having the first two domains of the human CD4 polypeptide and various domains of the constant

regions of the heavy or light chains of mammalian immunoglobulins have been described. (EP 394,827; Traunecker et al., 1988, *Nature*, 331:84-86). The polypeptides of the present invention fused or conjugated to an antibody, or portion thereof, having disulfide-linked dimeric structures (due to the IgG), for example, can also be more efficient in binding and neutralizing other molecules than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., 1995, *J. Biochem.*, 270:3958-3964). In many cases, the Fc portion in a fusion protein is beneficial in therapy, diagnosis, and/or screening methods, and thus can result in, for example, improved pharmacokinetic properties. (EP 232, 262 A). In drug discovery, for example, human proteins, such as huIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of huIL-5. (See, Bennett et al., 1995, *J. Molecular Recognition*, 8:52-58; and Johanson et al., 1995, *J. Biol. Chem.*, 270:9459-9471). Alternatively, deleting the Fc portion after the fusion protein has been expressed, detected, and purified may be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations.

Moreover, the antibodies or fragments thereof of the present invention can be fused to marker sequences, such as a peptide, to facilitate their purification. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., Chatsworth, CA), among others, many of which are commercially available. As described in Gentz et al., 1989, *Proc. Natl. Acad. Sci. USA*, 86:821-824, for instance, hexa-histidine provides for convenient purification of the fusion protein. Other peptide tags useful for purification include, but are not limited to, the "HA" tag, which corresponds to an epitope derived from the influenza hemagglutinin (HA) protein (Wilson et al., 1984, *Cell*, 37:767) and the "flag" tag.

The present invention further encompasses antibodies or fragments thereof conjugated to a diagnostic or therapeutic agent. The antibodies can be used diagnostically to, for example, monitor the development or progression of a tumor as part of a clinical testing procedure, for example, to determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling the antibody to a detectable substance. Nonlimiting examples of detectable substances include various

enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, and nonradioactive paramagnetic metal ions. The detectable substance can be coupled or conjugated either directly to the antibody
5 (or fragment thereof) or indirectly, through an intermediate (such as, for example, a linker known in the art) using techniques known in the art. (See, for example, U.S. Patent No. 4,741,900 for metal ions which can be conjugated to antibodies for use as diagnostics according to the present invention).

Nonlimiting examples of suitable enzymes include horseradish peroxidase,
10 alkaline phosphatase, beta-galactosidase, or acetylcholinesterase. Nonlimiting examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; nonlimiting examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; a nonlimiting
15 example of a luminescent material includes luminol; nonlimiting examples of bioluminescent materials include luciferase, luciferin, and aequorin; and nonlimiting examples of suitable radioactive material include iodine (^{125}I , ^{131}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium (^{111}In and other radioactive isotopes of indium), technetium (^{99}Tc , $^{99\text{m}}\text{Tc}$), thallium (^{201}Tl), gallium (^{68}Ga , ^{67}Ga), palladium (^{103}Pd),
20 molybdenum (^{99}Mo), xenon (^{133}Xe), fluorine (^{19}F), ^{153}Sm , ^{177}Lu , Gd, radioactive Pm, radioactive La, radioactive Yb, ^{166}Ho , ^{90}Y , radioactive Sc, radioactive Re, radioactive Re, ^{142}Pr , ^{105}Rh , and ^{97}Ru .

In specific embodiments, the protein tyrosine kinase biomarker polypeptides of the invention are attached to macrocyclic chelators useful for conjugating
25 radiometal ions, including, but not limited to, ^{111}In , ^{177}Lu , ^{90}Y , ^{166}Ho , and ^{153}Sm , to polypeptides. In a preferred embodiment, the radiometal ion associated with the macrocyclic chelators attached to the protein tyrosine kinase biomarker polypeptides of the invention is ^{111}In . In another preferred embodiment, the radiometal ion associated with the macrocyclic chelator attached to the protein tyrosine kinase
30 biomarker polypeptides of the invention is ^{90}Y . In specific embodiments, the macrocyclic chelator is 1, 4, 7, 10-tetraazacyclododecane-N, N', N'', N'''-tetraacetic

acid (DOTA). In other specific embodiments, the DOTA is attached to the protein tyrosine kinase biomarker polypeptides of the invention via a linker molecule.

Examples of linker molecules useful for conjugating DOTA to a polypeptide are commonly known in the art. (See, for example, DeNardo et al., 1998, *Clin. Cancer Res.*, 4(10):2483-90; Peterson et al., 1999, *Bioconjug. Chem.*, 10(4):553-557; and Zimmerman et al., 1999, *Nucl. Med. Biol.*, 26(8):943-950, which are hereby incorporated by reference in their entirety). In addition, U.S. Patent Nos. 5,652,361 and 5,756,065, which disclose chelating agents that can be conjugated to antibodies and methods for making and using them, are hereby incorporated by reference in their
10 entireties. Though U.S. Patent Nos. 5,652,361 and 5,756,065 focus on conjugating chelating agents to antibodies, one skilled in the art can readily adapt the methods disclosed therein in order to conjugate chelating agents to other polypeptides.

Antibodies can also be attached to solid supports, which are particularly useful for immunoassays or purification of the target antigen. Such solid supports include,
15 but are not limited to, glass, cellulose, polyacrylamide, nylon, polystyrene, polyvinyl chloride or polypropylene. Techniques for conjugating therapeutic moieties to antibodies are well known, see, e.g., Arnon et al., "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", In: *Monoclonal Antibodies And Cancer Therapy*, Reisfeld et al. (eds.), pp. 243-56, Alan R. Liss, Inc., 1985; Hellstrom et al., "Antibodies For Drug Delivery", In: *Controlled Drug Delivery* (2nd Ed.),
20 Robinson et al. (eds.), pp. 623-53, Marcel Dekker, Inc., 1987; Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", In: *Monoclonal Antibodies '84: Biological And Clinical Applications*, Pinchera et al. (eds.), pp. 475-506, 1985; "Analysis, Results, And Future Prospective Of The Therapeutic Use Of Radiolabeled Antibody In Cancer Therapy", In: *Monoclonal Antibodies For Cancer
25 Detection And Therapy*, Baldwin et al. (eds.), pp. 303-316, Academic Press, 1985; and Thorpe et al., 1982, "The Preparation And Cytotoxic Properties Of Antibody-Toxin Conjugates", *Immunol. Rev.*, 62:119-158. Alternatively, an antibody can be conjugated to a second antibody to form an antibody heteroconjugate, e.g., as
30 described in U.S. Patent No. 4,676,980 to Segal, which is incorporated herein by reference in its entirety. An antibody, i.e., an antibody specific for a protein tyrosine kinase biomarker polypeptide of this invention, with or without a therapeutic moiety

conjugated to it, and administered alone or in combination with cytotoxic factor(s) and/or cytokine(s), can be used as a therapeutic.

The antibodies of the invention can further be utilized for immunophenotyping of cell lines and biological samples. The translation product of the protein tyrosine kinase biomarker-encoding polynucleotides of the present invention can be useful as cell specific marker(s), or more specifically, as cellular marker(s) that are differentially expressed at various stages of differentiation and/or maturation of particular cell types. Monoclonal antibodies directed against a specific epitope, or combination of epitopes, allow for the screening of cellular populations expressing the marker. Various techniques utilizing monoclonal antibodies can be employed to screen for cellular populations expressing the marker(s), including magnetic separation using antibody-coated magnetic beads, "panning" with antibody(ies) attached to a solid matrix (i.e., tissue culture plate), and flow cytometry (See, e.g., U.S. Patent No. 5,985,660; Morrison et al., 1999, *Cell*, 96:737-749; and L.J. Wysocki and V.L. Sato, 1978, *Proc. Natl. Acad. Sci. USA*, 75(6):2844-8).

These techniques allow for the screening of particular populations of cells, such as might be found with hematological malignancies (i. e. minimal residual disease (MRD) in acute leukemic patients) and "non-self" cells in transplantations to prevent Graft-versus-Host Disease (GVHD). Alternatively, these techniques allow for the screening of hematopoietic stem and progenitor cells capable of undergoing proliferation and/or differentiation, as might be found in human umbilical cord blood.

Antibodies according to this invention can be assayed for immunospecific binding by any method known in the art. The immunoassays which can be used include, but are not limited to, competitive and non-competitive assay systems using techniques such as BIAcore analysis, FACS (Fluorescence Activated Cell Sorter) analysis, immunofluorescence, immunocytochemistry, Western blots, radioimmunoassays, ELISA (enzyme linked immunosorbent assays), "sandwich" immunoassays, immunoprecipitation assays, precipitin reactions, gel diffusion precipitin reactions, immunodiffusion assays, agglutination assays, complement fixation assays, immunoradiometric assays, fluorescent immunoassays, protein A immunoassays, to name but a few. Such assays are routine and well known and practiced in the art (see, e.g., Ausubel et al, eds, 1994, *Current Protocols in*

Molecular Biology, Vol. 1, John Wiley & Sons, Inc., New York, which is incorporated by reference herein in its entirety). Nonlimiting, exemplary immunoassays are described briefly below.

Immunoprecipitation protocols generally comprise lysing a population of cells
5 in a lysis buffer such as RIPA buffer (i.e., 1% NP-40 or Triton X-100, 1% sodium
deoxycholate, 0.1% SDS, 0.15 M NaCl, 0.01 M sodium phosphate at pH 7.2, 1%
Trasylol) supplemented with protein phosphatase and/or protease inhibitors (e.g.,
EDTA, PMSF, aprotinin, sodium vanadate); adding the antibody of interest to the cell
lysate; incubating for a period of time (e.g., 1 to 4 hours) at 4°C; adding protein A
10 and/or protein G sepharose beads to the cell lysate; incubating for about 60 minutes or
more at 4°C; washing the beads in lysis buffer; and resuspending the beads in
SDS/sample buffer. The ability of the antibody of interest to immunoprecipitate a
particular antigen can be assessed by, for example, Western blot analysis. One of skill
in the art would be knowledgeable as to the parameters that can be modified to
15 increase the binding of the antibody to an antigen and decrease the background (e.g.,
pre-clearing the cell lysate with sepharose beads). For further discussion regarding
immunoprecipitation protocols, see, e.g., Ausubel et al, eds, 1994, *Current Protocols
in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, at 10.16.1.

Western blot analysis generally comprises preparing protein samples;
20 electrophoresis of the protein samples in a polyacrylamide gel (e.g., 8%-20% SDS
PAGE depending on the molecular weight of the antigen); transferring the protein
sample from the polyacrylamide gel to a membrane such as nitrocellulose, PVDF or
nylon; blocking the membrane in blocking solution (e. g., PBS with 3% BSA or
nonfat milk); washing the membrane in washing buffer (e.g., PBS-Tween 20);
25 blocking the membrane with primary antibody (the antibody of interest) diluted in
blocking buffer; washing the membrane in washing buffer; blocking the membrane
with a secondary antibody (which recognizes the primary antibody, e.g., an anti-
human antibody) conjugated to an enzymatic substrate (e.g., horseradish peroxidase or
alkaline phosphatase) or radioactive molecule (e.g., ³²P or ¹²⁵I) diluted in blocking
30 buffer; washing the membrane in wash buffer; and detecting the presence of the
bound antigen. One of skill in the art would be knowledgeable as to the parameters
that can be modified to increase the signal detected and to reduce the background

noise. For further discussion regarding Western blot protocols, see, e.g., Ausubel et al, eds, 1994, *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, at 10.8.1.

5 ELISAs comprise preparing antigen; coating the wells of a 96 well microtiter plate with antigen; adding to the wells the antibody of interest conjugated to a detectable compound such as an enzymatic substrate (e.g., horseradish peroxidase or alkaline phosphatase); incubating for a period of time; and detecting the presence of the antigen. In ELISAs, the antibody of interest does not have to be conjugated to a detectable compound; instead, a second antibody (which recognizes the antibody of interest bound to antigen) conjugated to a detectable compound can be added to the wells. Further, instead of coating the wells with antigen, the antibodies can be first coated onto the well. In this case, a second antibody conjugated to a detectable compound can be added to the antibody-coated wells following the addition of the antigen of interest. One of skill in the art would be knowledgeable as to the parameters that can be modified to increase the signal detected, as well as other variations of ELISAs known in the art. For further discussion regarding ELISAs, see, e.g., Ausubel et al, eds, 1994, *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, at 11.2.1.

20 The binding affinity of an antibody to an antigen and the off-rate of an antibody-antigen interaction can be determined by competitive binding assays. One example of a competitive binding assay is a radioimmunoassay involving the incubation of labeled antigen (e.g., ^3H or ^{125}I), or a fragment or variant thereof, with the antibody of interest in the presence of increasing amounts of labeled antigen, and the detection of the antibody bound to the labeled antigen. The affinity of the antibody of interest for a protein tyrosine kinase biomarker and the binding off rates can be determined from the data by Scatchard plot analysis. Competition with a second antibody can also be determined using radioimmunoassays. In this case, the tyrosine kinase biomarker protein is incubated with an antibody of interest conjugated to a labeled compound (e.g., a compound labeled with ^3H or ^{125}I) in the presence of increasing amounts of an unlabeled second antibody. This kind of competitive assay between two antibodies can also be used to determine if two antibodies bind to the same or different epitopes on an antigen.

In a preferred embodiment, BIAcore kinetic analysis is used to determine the binding on and off rates of antibodies (including antibody fragments or variants thereof) to a tyrosine kinase biomarker protein, or fragments of a tyrosine kinase biomarker protein. Kinetic analysis comprises analyzing the binding and dissociation of antibodies from chips with immobilized tyrosine kinase biomarker protein on the chip surface.

It is to be further understood that the above-described techniques for the production, expression, isolation, and manipulation of antibody molecules, for example, by recombinant techniques involving molecular biology, as well as by other techniques related to the analysis of polynucleotides and proteins, are applicable to other polypeptide or peptide molecules of the invention as described herein, in particular, the tyrosine kinase biomarker polypeptides or peptides themselves, as applicable or warranted. in accordance with the various embodiments of this invention.

The present invention also embraces a kit for determining, predicting, or prognosing drug susceptibility or resistance by a patient having a disease, particularly a cancer or tumor, preferably, a breast cancer or tumor. Such kits are useful in a clinical setting for use in testing patient's biopsied tumor or cancer samples, for example, to determine or predict if the patient's tumor or cancer will be resistant or sensitive to a given treatment or therapy with a drug, compound, chemotherapy agent, or biological treatment agent. Provided in the kit are the predictor set comprising those polynucleotides correlating with resistance and sensitivity to protein tyrosine kinase modulators in a particular biological system, particularly protein tyrosine kinase inhibitors, and preferably comprising a microarray; and, in suitable containers, the modulator compounds for use in testing cells from patient tissue or patient samples for resistance/sensitivity; and instructions for use. Such kits encompass predictor set comprising those polynucleotides correlating with resistance and sensitivity to modulators of protein tyrosine kinases including members of the Src family of tyrosine kinases, for example, Src, Fgr, Fyn, Yes, Blk, Hck, Lck and Lyn, as well as other protein tyrosine kinases, including, Bcr-abl, Jak, PDGFR, c-kit and Eph receptors,

Also, as explained above, the kit is not limited to microarrays, but can encompass a variety of methods and systems by which the expression of the predictor/marker polynucleotides can be assayed and/or monitored, both at the level of mRNA and of protein, for example, via PCR assays, e.g., RT-PCR and immunoassay, such as ELISA. In kits for performing PCR, or *in situ* hybridization, for example, nucleic acid primers or probes from the sequences of one or more of the predictor polynucleotides, such as those described herein, in addition to buffers and reagents as necessary for performing the method, and instructions for use. In kits for performing immunoassays, e.g. ELISAs, immunoblotting assays, and the like, antibodies, or bindable portions thereof, to the protein tyrosine kinase biomarker polypeptides of the invention, or to antigenic or immunogenic peptides thereof, are supplied, in addition to buffers and reagents as necessary for performing the method, and instructions for use. The kits according to the present invention can also comprise predictor polynucleotides as set forth in Table 2, and/or one or more of the specific predictor polynucleotide subsets as presented in Tables 4-5 herein.

In another embodiment, the present invention embraces the use of one or more polynucleotides among those of the predictor polynucleotides identified herein that can serve as targets for the development of drug therapies for disease treatment. Such targets may be particularly applicable to treatment of breast diseases, such as breast cancers or tumors. Indeed, because these predictor polynucleotides are differently expressed in sensitive and resistant cells, their expression pattern is correlated with relative intrinsic sensitivity of cells to treatment with compounds that interact with and inhibit protein tyrosine kinases. Accordingly, the polynucleotides highly expressed in resistant cells can serve as targets for the development of drug therapies for the tumors which are resistant to protein tyrosine kinase inhibitor compounds, for example, Src tyrosine kinase inhibitors.

In another embodiment, the present invention embraces antisense and/or siRNAi reagents as specific modulators of the predictor polynucleotides of the present invention. In specific embodiments, antagonists according to the present invention are nucleic acids corresponding to the sequences contained in one or more of the sequences provided as SEQ ID NO:1 thru 137, or the complementary strand thereof. In one embodiment, antisense sequence is generated internally by the organism, in

another embodiment, the antisense sequence is separately administered (see, for example, O'Connor, *Neurochem.*, 56:560 (1991). *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988). Antisense technology can be used to control gene expression through antisense DNA or RNA, or through triple-helix formation. Antisense techniques are discussed for example, in Okano, *Neurochem.*, 56:560 (1991); *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance, Lee et al., *Nucleic Acids Research*, 6:3073 (1979); Cooney et al., *Science*, 241:456 (1988); and Dervan et al., *Science*, 251:1300 (1991). The methods are based on binding of a polynucleotide to a complementary DNA or RNA.

For example, the use of c-myc and c-myb antisense RNA constructs to inhibit the growth of the non-lymphocytic leukemia cell line HL-60 and other cell lines was previously described. (Wickstrom et al. (1988); Anfossi et al. (1989)). These experiments were performed in vitro by incubating cells with the oligoribonucleotide. A similar procedure for in vivo use is described in WO 91/15580. Briefly, a pair of oligonucleotides for a given antisense RNA is produced as follows: A sequence complimentary to the first 15 bases of the open reading frame is flanked by an EcoRI site on the 5' end and a HindIII site on the 3' end. Next, the pair of oligonucleotides is heated at 90°C for one minute and then annealed in 2X ligation buffer (20mM TRIS HCl pH 7.5, 10mM MgCl₂, 10mM dithiothreitol (DTT) and 0.2 mM ATP) and then ligated to the EcoRI/Hind III site of the retroviral vector PMV7 (WO 91/15580).

For example, the 5' coding portion of a polynucleotide that encodes the mature polypeptide of the present invention may be used to design an antisense RNA oligonucleotide of from about 10 to 40 base pairs in length. A DNA oligonucleotide is designed to be complementary to a region of the gene involved in transcription thereby preventing transcription and the production of the receptor. The antisense RNA oligonucleotide hybridizes to the mRNA in vivo and blocks translation of the mRNA molecule into receptor polypeptide. Antisense oligonucleotides may be single or double stranded. Double stranded RNA's may be designed based upon the teachings of Paddison et al., *Proc. Nat. Acad. Sci.*, 99:1443-1448 (2002); and International Publication Nos. WO 01/29058, and WO 99/32619; which are hereby incorporated herein by reference.

In one embodiment, the antisense nucleic acid of the invention is produced intracellularly by transcription from an exogenous sequence. For example, a vector or a portion thereof, is transcribed, producing an antisense nucleic acid (RNA) of the invention. Such a vector would contain a sequence encoding the antisense nucleic acid of the invention. Such a vector can remain episomal or become chromosomally integrated, as long as it can be transcribed to produce the desired antisense RNA. Such vectors can be constructed by recombinant DNA technology methods standard in the art. Vectors can be plasmid, viral, or others known in the art, used for replication and expression in vertebrate cells. Expression of the sequence encoding a polypeptide of the invention, or fragments thereof, can be by any promoter known in the art to act in vertebrate, preferably human cells. Such promoters can be inducible or constitutive. Such promoters include, but are not limited to, the SV40 early promoter region (Bernoist and Chambon, *Nature*, 29:304-310 (1981), the promoter contained in the 3' long terminal repeat of Rous sarcoma virus (Yamamoto et al., *Cell*, 22:787-797 (1980), the herpes thymidine promoter (Wagner et al., *Proc. Natl. Acad. Sci. U.S.A.*, 78:1441-1445 (1981), the regulatory sequences of the metallothionein gene (Brinster et al., *Nature*, 296:39-42 (1982)), etc.

The antisense nucleic acids of the invention comprise a sequence complementary to at least a portion of an RNA transcript of a gene of interest. However, absolute complementarity, although preferred, is not required. A sequence "complementary to at least a portion of an RNA" referred to herein, means a sequence having sufficient complementarity to be able to hybridize with the RNA, forming a stable duplex; in the case of double stranded antisense nucleic acids of the invention, a single strand of the duplex DNA may thus be tested, or triplex formation may be assayed. The ability to hybridize will depend on both the degree of complementarity and the length of the antisense nucleic acid. Generally, the larger the hybridizing nucleic acid, the more base mismatches with a RNA sequence of the invention it may contain and still form a stable duplex (or triplex as the case may be). One skilled in the art can ascertain a tolerable degree of mismatch by use of standard procedures to determine the melting point of the hybridized complex.

Oligonucleotides that are complementary to the 5' end of the message, e.g., the 5' untranslated sequence up to and including the AUG initiation codon, should work

most efficiently at inhibiting translation. However, sequences complementary to the 3' untranslated sequences of mRNAs have been shown to be effective at inhibiting translation of mRNAs as well. See generally, Wagner, R., *Nature*, 372:333-335 (1994). Thus, oligonucleotides complementary to either the 5' - or 3' - non-translated, non-coding regions of a polynucleotide sequence of the invention could be used in an antisense approach to inhibit translation of endogenous mRNA. Oligonucleotides complementary to the 5' untranslated region of the mRNA should include the complement of the AUG start codon. Antisense oligonucleotides complementary to mRNA coding regions are less efficient inhibitors of translation but could be used in accordance with the invention. Whether designed to hybridize to the 5' -, 3' - or coding region of mRNA, antisense nucleic acids should be at least six nucleotides in length, and are preferably oligonucleotides ranging from 6 to about 50 nucleotides in length. In specific aspects the oligonucleotide is at least 10 nucleotides, at least 17 nucleotides, at least 25 nucleotides or at least 50 nucleotides.

The polynucleotides of the invention can be DNA or RNA or chimeric mixtures or derivatives or modified versions thereof, single-stranded or double-stranded. The oligonucleotide can be modified at the base moiety, sugar moiety, or phosphate backbone, for example, to improve stability of the molecule, hybridization, etc. The oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors in vivo), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., *Proc. Natl. Acad. Sci. U.S.A.* 86:6553-6556 (1989); Lemaitre et al., *Proc. Natl. Acad. Sci.*, 84:648-652 (1987); PCT Publication NO: WO88/09810, published December 15, 1988) or the blood-brain barrier (see, e.g., PCT Publication NO: WO89/10134, published April 25, 1988), hybridization-triggered cleavage agents. (See, e.g., Krol et al., *BioTechniques*, 6:958-976 (1988)) or intercalating agents. (See, e.g., Zon, *Pharm. Res.*, 5:539-549 (1988)). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered cleavage agent, etc.

Double stranded RNA's may be designed based upon the teachings of Paddison et al., *Proc. Nat. Acad. Sci.*, 99:1443-1448 (2002); and International

Publication Nos. WO 01/29058, and WO 99/32619; which are hereby incorporated herein by reference.

SiRNA reagents are specifically contemplated by the present invention. Such reagents are useful for inhibiting expression of the polynucleotides of the present invention and may have therapeutic efficacy. Several methods are known in the art
5 for the therapeutic treatment of disorders by the administration of siRNA reagents. One such method is described by Tiscornia et al (PNAS, 100(4):1844-1848 (2003)), which is incorporated by reference herein in its entirety.

The antisense oligonucleotide may comprise at least one modified base moiety
10 which is selected from the group including, but not limited to, 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-
15 methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-
20 thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine.

The antisense oligonucleotide may also comprise at least one modified sugar moiety selected from the group including, but not limited to, arabinose, 2-
25 fluoroarabinose, xylulose, and hexose.

In yet another embodiment, the antisense oligonucleotide comprises at least one modified phosphate backbone selected from the group including, but not limited to, a phosphorothioate, a phosphorodithioate, a phosphoramidothioate, a phosphoramidate, a phosphordiamidate, a methylphosphonate, an alkyl
30 phosphotriester, and a formacetal or analog thereof.

In yet another embodiment, the antisense oligonucleotide is an a-anomeric oligonucleotide. An a-anomeric oligonucleotide forms specific double-stranded

hybrids with complementary RNA in which, contrary to the usual b-units, the strands run parallel to each other (Gautier et al., Nucl. Acids Res., 15:6625-6641 (1987)). The oligonucleotide is a 2-O-methylribonucleotide (Inoue et al., Nucl. Acids Res., 15:6131-6148 (1987)), or a chimeric RNA-DNA analogue (Inoue et al., FEBS Lett. 5 215:327-330 (1987)).

Polynucleotides of the invention may be synthesized by standard methods known in the art, e.g. by use of an automated DNA synthesizer (such as are commercially available from Biosearch, Applied Biosystems, etc.). As examples, phosphorothioate oligonucleotides may be synthesized by the method of Stein et al. 10 (Nucl. Acids Res., 16:3209 (1988)), methylphosphonate oligonucleotides can be prepared by use of controlled pore glass polymer supports (Sarin et al., Proc. Natl. Acad. Sci. U.S.A., 85:7448-7451 (1988)), etc.

While antisense nucleotides complementary to the coding region sequence of the invention could be used, those complementary to the transcribed untranslated 15 region are most preferred.

Potential antagonists according to the invention also include catalytic RNA, or a ribozyme (See, e.g., PCT International Publication WO 90/11364, published October 4, 1990; Sarver et al, Science, 247:1222-1225 (1990). While ribozymes that cleave mRNA at site specific recognition sequences can be used to destroy mRNAs 20 corresponding to the polynucleotides of the invention, the use of hammerhead ribozymes is preferred. Hammerhead ribozymes cleave mRNAs at locations dictated by flanking regions that form complementary base pairs with the target mRNA. The sole requirement is that the target mRNA have the following sequence of two bases: 5'-UG-3'. The construction and production of hammerhead ribozymes is well known 25 in the art and is described more fully in Haseloff and Gerlach, Nature, 334:585-591 (1988). There are numerous potential hammerhead ribozyme cleavage sites within each nucleotide sequence disclosed in the sequence listing. Preferably, the ribozyme is engineered so that the cleavage recognition site is located near the 5' end of the mRNA corresponding to the polynucleotides of the invention; i.e., to increase 30 efficiency and minimize the intracellular accumulation of non-functional mRNA transcripts.

As in the antisense approach, the ribozymes of the invention can be composed of modified oligonucleotides (e.g. for improved stability, targeting, etc.) and should be delivered to cells which express the polynucleotides of the invention in vivo. DNA constructs encoding the ribozyme may be introduced into the cell in the same manner as described above for the introduction of antisense encoding DNA. A preferred method of delivery involves using a DNA construct "encoding" the ribozyme under the control of a strong constitutive promoter, such as, for example, pol III or pol II promoter, so that transfected cells will produce sufficient quantities of the ribozyme to destroy endogenous messages and inhibit translation. Since ribozymes unlike antisense molecules, are catalytic, a lower intracellular concentration is required for efficiency.

Antagonist/agonist compounds may be employed to inhibit the cell growth and proliferation effects of the polypeptides of the present invention on neoplastic cells and tissues, i.e. stimulation of angiogenesis of tumors, and, therefore, retard or prevent abnormal cellular growth and proliferation, for example, in tumor formation or growth.

The antagonist/agonist may also be employed to prevent hyper-vascular diseases, and prevent the proliferation of epithelial lens cells after extracapsular cataract surgery. Prevention of the mitogenic activity of the polypeptides of the present invention may also be desirous in cases such as restenosis after balloon angioplasty.

The antagonist/agonist may also be employed to prevent the growth of scar tissue during wound healing.

The antagonist/agonist may also be employed to treat, prevent, and/or diagnose the diseases described herein.

Thus, the invention provides a method of treating or preventing diseases, disorders, and/or conditions, including but not limited to the diseases, disorders, and/or conditions listed throughout this application, associated with overexpression of a polynucleotide of the present invention by administering to a patient (a) an antisense molecule directed to the polynucleotide of the present invention, and/or (b) a ribozyme directed to the polynucleotide of the present invention.

EXAMPLES

The Examples herein are meant to exemplify the various aspects of carrying out the invention and are not intended to limit the scope of the invention in any way. The Examples do not include detailed descriptions for conventional methods employed, such as in the construction of vectors, the insertion of cDNA into such vectors, or the introduction of the resulting vectors into the appropriate host. Such methods are well known to those skilled in the art and are described in numerous publications, for example, Sambrook, Fritsch, and Maniatis, Molecular Cloning: A Laboratory Manual, 2nd Edition, Cold Spring Harbor Laboratory Press, USA, (1989).

10

EXAMPLE 1 – METHODS

IC₅₀ determination--*in vitro* cytotoxicity assay

The protein tyrosine kinase inhibitor compound (described in international application WO 00/62778, published October 26, 2000) was tested for cytotoxicity *in vitro* against a panel of twenty-three human breast cell lines available from the American Type Culture Collection, ATCC, except H3396, which was obtained from Pacific Northwest Research Institute, Seattle WA. The MCF7/Her2 cell line was established by stable transfection of MCF7 cells with the HER2 gene. Cytotoxicity was assessed in cells by the MTS (3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxy-methoxyphenyl)-2-(4-sulphenyl)-2H-tetrazolium, inner salt) assay (T.L. Riss et al., 1992, *Mol. Biol. Cell*, 3 (Suppl.):184a).

To carry out the assays, the breast cells were plated at 4,000 cells/well in 96 well microtiter plates, and 24 hours later, serially diluted drugs were added. The concentration range for the protein tyrosine kinase inhibitor compound BMS-A used in the cytotoxicity assay was from 5 µg/ml to 0.0016 µg/ml (roughly 10 µM to 0.0032 µM).

The cells were incubated at 37°C for 72 hours at which time the tetrazolium dye, MTS (333 µg/ml final concentration), in combination with the electron coupling agent phenazine methosulfate (25 µM final concentration), was added. A dehydrogenase enzyme in live cells reduces the MTS to a form that absorbs light and can be quantified spectrophotometrically at 492 nM. The greater the absorbency the

30

greater the number of live cells. The results are expressed as an IC_{50} , which is the drug concentration required to inhibit cell proliferation (i.e. absorbance at 492 nm) to 50% of that of untreated control cells. The mean IC_{50} and standard deviation (SD) from multiple tests for each cell line were calculated.

5

Resistant/sensitivity classification

The IC_{50} of the BMS-A protein tyrosine kinase inhibitor compound for each cell line was log-transformed to $\log_{10}(IC_{50})$, and the mean $\log_{10}(IC_{50})$ across the 23 human breast cell lines was calculated. The resistance/sensitivity phenotype of the cell lines was classified as follows: the cell lines with $\log_{10}(IC_{50})$ below the mean
10 $\log_{10}(IC_{50})$ of all 23 cell lines were defined as sensitive to the compound, while those with $\log_{10}(IC_{50})$ above the mean $\log_{10}(IC_{50})$ were considered to be resistant to the compound. The resistance/sensitivity classification is shown in Table 1 and 7 cell lines classified as sensitive and 16 cell lines classified as resistant to the protein tyrosine kinase inhibitor compound BMS-A.

15

Polynucleotide expression profiling

The breast cells were grown under standard cell culture conditions: RPMI 1640 supplemented to contain 10% fetal bovine serum, 100 IU/ml penicillin, 100 mg/ml streptomycin, 2 mM L-glutamine and 10 mM Hepes (all from GibcoBRL, Rockville, MD). RNA was isolated from the cultured cells, either treated or untreated
20 with drug (i.e., the protein tyrosine kinase inhibitor compound) at 50-70% confluence using the RNeasy™ kits commercially available from Qiagen, Valencia, CA. The quality of the RNA was assessed by measuring the 28s:18s ribosomal RNA ratio using an Agilent 2100 bioanalyzer (Agilent Technologies, Rockville, MD). The concentration of total RNA was determined spectrophotometrically. 10 µg of total
25 RNA from each cell line was used to prepare biotinylated probe according to the Affymetrix Genechip® Expression Analysis Technical Manual, 2001. Targets were hybridized to Affymetrix high density oligonucleotide array human HG-U133 set chips (Affymetrix, Santa Clara, CA). The arrays were then washed and stained using the GeneChip® Fluidics station according to the manufacture's instructions
30 (Affymetrix Genechip® Technical Manual, 2001). The HG-U133 set contains 2 Genechip® arrays, which contain approximately 45,000 probe sets representing more

than 39,000 transcripts derived from approximately 33,000 well-substantiated human polynucleotides.

Preprocessing of microarray data

Scanned image files were visually inspected for artifacts and analyzed with
5 GeneChip® Expression Analysis software MAS 5.0 (Affymetrix, Santa Clara, CA).
The "Detection Call" (Affymetrix Genechip® Expression Analysis Technical Manual,
2001) is used to determine whether a transcript is detected within one sample; the
"Signal" (Affymetrix Genechip® Expression Analysis Technical Manual, 2001)
measures the relative abundance of a transcript. The trimmed mean intensity for each
10 chip was scaled to 1,500 (see, Affymetrix Genechip® Expression Analysis Technical
Manual, 2001) in order to account for any minor differences in global chip intensity,
so that the overall expression level for each cell line was comparable. Affymetrix
control sequences were removed prior to analysis.

Of a total of 44,792 probe sets on the HG-U133 arrays, 15,707 represented
15 probe sets were not detected (Absent Call; p-value >0.06) across all of the 23 breast
cell lines using the Affymetrix GeneChip® Expression Analysis algorithm; these
undetected polynucleotides were excluded from further analysis.

The remaining data containing 29,085 probe sets were transferred to the
GeneCluster software (Whitehead Institute; T.R. Golub et al., 1999, *Science*, 286:531-
20 537). A threshold filter was applied to the polynucleotide expression values of the
remaining data to remove low and high polynucleotide expression values that were
not likely to be in the linear range of the Affymetrix fluorescent scanner. The
threshold filter converted all polynucleotide expression values that were below 100
units to 100 units, and all polynucleotide expression values that were above 45,000
25 units to 45,000 units. All represented polynucleotides whose polynucleotide
expression values were between 100 and 45,000 were not changed.

A second "variation filter" was then applied to the data set to find
polynucleotides that were likely to correlate with different properties and features of
the 23 cell lines. The object of the second filter was to select those polynucleotides
30 whose expression pattern varied across the data set, because a polynucleotide that
does not vary can not provide information about differing properties of the 23 cell line

panel. For example, if there are two populations of cells within the data set, e.g., fast growing cells and slow growing cells, then a polynucleotide whose expression is constant, or whose expression does not change substantially, can not yield information that would correlate to fast or slow cell growth.

5 The second variation filter was formulated to determine the expression pattern of each polynucleotide across the 23 breast cell lines and to find polynucleotides that passed the following criteria:

1. The polynucleotide must show a three-fold change in absolute expression, i.e., as depicted in the formula:

10

$$\frac{\text{expression value in any given cell line}}{\text{expression value in any other cell line}} > 3 \text{ or } < 0.33$$

15 2. In addition to 1, the three-fold change must represent an absolute difference of 1000 expression units.

3. In addition, the criteria in #1 and #2 above must be met on four independent occasions within the data set, i.e., Cell line A/B, Cell line E/F, Cell line C/U and Cell line T/G. (The algorithm does not use a single expression value for one cell line on multiple occasions, i.e., Cell Line A/B, Cell line A/G, Cell line A/F and
20 Cell line B/F).

The second variation filter reduced the data set to 5322 polynucleotides. After the second variation filter, the expression value for each polynucleotide was log transformed and normalized to the mean across all of the 23 samples (mean set to 0 and standard deviation set to 1). This normalized data set was used to select
25 polynucleotides which significantly correlated with the property of sensitivity toward a drug class as described herein.

Drug (BMS-A) treatment of breast cell lines and selection of
polynucleotides modified by the drug

30 The 11 breast cell lines (indicated in bold in the Table 1) with an IC₅₀ ranging from 0.0055 to 9.5 µM were used in a drug induction study employing the BMS-A protein tyrosine kinase inhibitor. Cells were seeded in a 10 cm² culture plate in cell culture medium as described herein and were cultured for 24 hours at 37°C. The

medium was then changed to medium containing drug (0.4 μ M BMS-A compound in 0.1% DMSO, Sigma); the cells were incubated for another 24 hours, and then lysed for RNA isolation. The expression profiling was performed as described above and data were analyzed using GeneChip[®] Expression Analysis software MAS 5.0
5 (Affymetrix, Santa Clara, CA). The expression data of a drug treated cell line were compared pair-wise to data from the same cell line untreated with drug. A change in p-value was calculated, indicating an increase, decrease or no change in polynucleotide expression. When the p-value was less than 0.0025, the change was considered to be significant. This analysis was performed for all 11 cell lines to
10 compare the polynucleotide expression with or without drug treatment.

EXAMPLE 2 – PCR EXPRESSION PROFILING

RNA quantification is performed using the Taqman[®] real-time-PCR fluorogenic assay. The Taqman[®] assay is one of the most precise methods for
15 assaying the concentration of nucleic acid templates.

RNA is prepared using standard methods, preferably, employing the RNeasy Maxi Kit commercially available from Qiagen (Valencia, CA). A cDNA template for real-time PCR can be generated using the Superscript[™] First Strand Synthesis system for RT-PCR. Representative forward and reverse RT-PCT primers for each of the
20 protein tyrosine kinase biomarker polynucleotides of the present invention are provided in Table 6.

SYBR Green real-time PCR reactions are prepared as follows: The reaction mix contains 20 ng first strand cDNA; 50 nM Forward Primer; 50 nM Reverse Primer; 0.75X SYBR Green I (Sigma); 1X SYBR Green PCR Buffer (50 mM Tris-HCl pH 8.3, 75 mM KCl); 10% DMSO; 3 mM MgCl₂; 300 μ M each dATP, dGTP, dTTP, dCTP; 1 U Platinum[®] Taq DNA Polymerase High Fidelity (Cat# 11304-029; Life Technologies; Rockville, MD). Real-time PCR is performed using an Applied Biosystems 5700 Sequence Detection System. Conditions are 95°C for 10 minutes (denaturation and activation of Platinum[®] Taq DNA Polymerase), 40 cycles of PCR
25 (95°C for 15 seconds, 60°C for 1 minute). PCR products are analyzed for uniform
30 melting using an analysis algorithm built into the 5700 Sequence Detection System.

cDNA quantification used in the normalization of template quantity is performed using Taqman® technology. Taqman® reactions are prepared as follows: The reaction mix comprises 20 ng first strand cDNA; 25 nM GAPDH-F3, Forward Primer; 250 nM GAPDH-R1 Reverse Primer; 200 nM GAPDH-PVIC Taqman®
 5 Probe (fluorescent dye labeled oligonucleotide primer); 1X Buffer A (Applied Biosystems); 5.5 mM MgCl₂; 300 μM dATP, dGTP, dTTP, dCTP; and 1 U Amplitaq Gold (Applied Biosystems). GAPDH (D-glyceraldehyde-3-phosphate dehydrogenase) is used as a control to normalize mRNA levels. Real-time Taqman® PCR is performed using an Applied Biosystems 7700 Sequence Detection System.
 10 Conditions are 95°C for 10 minutes (denaturation and activation of Amplitaq Gold), 40 cycles of PCR (95°C for 15 seconds, 60°C for 1 minute).

The sequences for the GAPDH oligonucleotides used in the Taqman® reactions are as follows:

- 15 GAPDH-F3: 5'-AGCCGAGCCACATCGCT-3' (SEQ ID NO:531);
 GAPDH-R1: 5'-GTGACCAGGCGCCCAATAC-3' (SEQ ID NO:532); and
 GAPDH-PVIC Taqman® Probe -VIC-5'-
 CAAATCCGTTGACTCCGACCTTCACCTT-3' TAMRA (SEQ ID NO:533).

20 The Sequence Detection System generates a Ct (threshold cycle) value that is used to calculate a concentration for each input cDNA template. cDNA levels for each polynucleotide of interest are normalized to GAPDH cDNA levels to compensate for variations in total cDNA quantity in the input sample. This is done by generating GAPDH Ct values for each cell line. Ct values for the polynucleotide of
 25 interest and GAPDH are inserted into a modified version of the $\delta\delta Ct$ equation (Applied Biosystems Prism® 7700 Sequence Detection System User Bulletin #2), which is used to calculate a GAPDH normalized relative cDNA level for each specific cDNA. The $\delta\delta Ct$ equation is as follows: relative quantity of nucleic acid template
 $= 2^{\delta\delta Ct} = 2^{(\delta Ct_a - \delta Ct_b)}$, where $\delta Ct_a = Ct_{\text{target}} - Ct_{\text{GAPDH}}$, and $\delta Ct_b = Ct_{\text{reference}} - Ct_{\text{GAPDH}}$. (No reference cell line is used for the calculation of relative quantity; δCt_b
 30 is defined as 21).

EXAMPLE 3 – PRODUCTION OF AN ANTIBODY DIRECTED AGAINST PROTEIN TYROSINE KINASE BIOMARKER POLYPEPTIDES

Anti-protein tyrosine kinase biomarker polypeptide antibodies of the present
5 invention can be prepared by a variety of methods as detailed hereinabove. As one
example of an antibody-production method, cells expressing a polypeptide of the
present invention are administered to an animal as immunogen to induce the
production of sera containing polyclonal antibodies directed against the expressed
polypeptide. In a preferred method, the expressed polypeptide is prepared, preferably
10 isolated and/or purified, to render it substantially free of natural contaminants using
techniques commonly practiced in the art. Such a preparation is then introduced into
an animal in order to produce polyclonal antisera of greater specific activity for the
expressed and isolated polypeptide.

In a most preferred method, the antibodies of the present invention are
15 monoclonal antibodies (or protein binding fragments thereof) and can be prepared
using hybridoma technology as detailed hereinabove. Cells expressing the
polypeptide can be cultured in any suitable tissue culture medium; however, it is
frequently preferable to culture cells in Earle's modified Eagle's medium
supplemented to contain 10% fetal bovine serum (inactivated at about 56°C), and
20 supplemented to contain about 10 g/l nonessential amino acids, about 1.0 U/ml
penicillin, and about 100 µg/ml streptomycin.

The splenocytes of immunized (and boosted) mice are extracted and fused
with a suitable myeloma cell line. Any suitable myeloma cell line can be employed in
accordance with the present invention; however, it is preferable to employ the parent
25 myeloma cell line SP2/0, available from the ATCC. After fusion, the resulting
hybridoma cells are selectively maintained in HAT medium, and then cloned by
limiting dilution as described, for example, by Wands et al. (1981, *Gastroenterology*,
80:225-232). The hybridoma cells obtained through such a selection process are then
assayed to identify those cell clones that secrete antibodies capable of binding to the
30 polypeptide immunogen, or a portion thereof.

Alternatively, additional antibodies capable of binding to the polypeptide can
be produced in a two-step procedure using anti-idiotypic antibodies. Such a method

makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain a second antibody that binds to a first antibody. In accordance with this method, protein-specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an immunized animal are then used to produce
5 hybridoma cells, and the hybridoma cells are screened to identify clones that produce an antibody whose ability to bind to the protein-specific antibodies can be blocked by the protein. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and can be used to immunize an animal to induce the formation of further protein-specific antibodies.

10 For *in vivo* use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known and practiced in the art. (See, e.g., for review, Morrison, 1985, *Science*, 229:1202); Oi et al., 1986,
15 *BioTechniques*, 4:214; Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., 1984, *Nature*, 312:643; and Neuberger et al., 1985, *Nature*, 314:268).

20 EXAMPLE 4 – IMMUNOFLUORESCENCE ASSAYS

The following immunofluorescence protocol can be used, for example, to verify protein tyrosine kinase biomarker expression in cells, or, for example, to check for the presence of one or more antibodies that bind protein tyrosine kinase biomarkers (polypeptides or peptides) expressed on the surfaces of cells. Briefly,
25 Lab-Tek II chamber slides are coated overnight at 4°C with 10 µg/ml of bovine collagen Type II in DPBS containing calcium and magnesium (DPBS++). The slides are then washed twice with cold DPBS++ and seeded with approximately 8000 CHO cells transfected with a vector comprising the coding sequence for a protein tyrosine kinase biomarker of the present invention or with CHO cells transfected with vector
30 alone (control) in a total volume of 125 µl and incubated at 37°C in the presence of 95% oxygen / 5% carbon dioxide.

Thereafter, the culture medium is gently removed by aspiration and the adherent cells are washed twice with DPBS++ at ambient temperature. The slides are blocked with DPBS++ containing 0.2% BSA (blocker) at 0-4°C for one hour. The blocking solution is gently removed by aspiration, and 125 µl of antibody containing
5 solution (an antibody containing solution may be, for example, a hybridoma culture supernatant which is usually used undiluted, or serum/plasma which is usually diluted, e.g., a dilution of about 1:50, 1:100, 1:1000, and the like). The slides are incubated for 1 hour at 0-4°C. Antibody solutions are then gently removed by aspiration and the cells are washed 5 times with 400 µl of ice cold blocking solution. Next, 125 µl of 1
10 µg/ml rhodamine labeled secondary antibody (e.g., anti-human IgG) in blocker solution is added to the cells. Again, cells are incubated for 1 hour at 0-4°C.

The secondary antibody solution is then gently removed by aspiration and the cells are washed 3 times with 400 µl of ice cold blocking solution, and 5 times with cold DPBS++. The cells are then fixed with 125 µl of 3.7% formaldehyde in
15 DPBS++ for 15 minutes at ambient temperature. Thereafter, the cells are washed 5 times with 400 µl of DPBS++ at ambient temperature. Finally, the cells are mounted in 50% aqueous glycerol and viewed using a fluorescence microscope using rhodamine filters.

20 EXAMPLE 5 – COMPLIMENTARY SEQUENCES.

Antisense molecules or nucleic acid sequences complementary to the protein tyrosine kinase biomarker polypeptides-encoding sequence, or any part thereof, is used to decrease or to inhibit the expression of naturally occurring protein tyrosine kinase biomarker polypeptides. Although the use of antisense or complementary
25 oligonucleotides comprising about 15 to 35 base-pairs is described, essentially the same procedure is used with smaller or larger nucleic acid sequence fragments. An oligonucleotide based on the coding sequence of protein tyrosine kinase biomarker polypeptides, as depicted in SEQ ID NO:1 thru 137, for example, is used to inhibit expression of naturally occurring protein tyrosine kinase biomarker polypeptides. The
30 complementary oligonucleotide is typically designed from the most unique 5' sequence and is used either to inhibit transcription by preventing promoter binding to

the coding sequence, or to inhibit translation by preventing the ribosome from binding to the protein tyrosine kinase biomarker polypeptides-encoding transcript, among others. However, other regions may also be targeted.

Using an appropriate portion of the signal and 5' sequence of SEQ ID NO:1 thru 137, an effective antisense oligonucleotide includes any of about 15-35 nucleotides spanning the region which translates into the signal or 5' coding sequence, among other regions, of the polypeptide as depicted in SEQ ID NO:138 thru 256. Appropriate oligonucleotides may be designed using OLIGO 4.06 software and the protein tyrosine kinase biomarker polypeptides coding sequence (SEQ ID NO:1 thru 137). Preferred oligonucleotides are deoxynucleotide, or chimeric deoxynucleotide/ribonucleotide based and are provided below. The oligonucleotides may be synthesized using chemistry essentially as described in U.S. Patent No. 5,849,902; which is hereby incorporated herein by reference in its entirety.

Representative RNAi reagent sequences are as follows:

Target Name	Sense Strand RNAi Reagent	SEQ ID NO:	Anti-Sense Strand RNAi Reagent	SEQ ID NO:
caveolin 1-1	CAGGGCAACAUCUACAA GCTT	534	GCUUGUAGAUGUUGCCCU GTT	546
caveolin 1-2	GCAAGUGUACGACGCGC ACTT	535	GUGCGCGUCGUACACUUG CTT	547
caveolin 1-3	CCGCUUGCUGUCUGCCCU CTT	536	GAGGGCAGACAGCAAGCG GTT	548
caveolin 1-4	CAUCUGGGCAGUUGUAC CATT	537	UGGUACAACUGCCCAGAU GTT	549
caveolin 2-1	CUACGCACUCCUUUGACA ATT	538	UUGUCAAAAGGAGUGCGUA GTT	550
caveolin 2-2	AGUGUGGAUCUGCAGCC AUTT	539	AUGGCUGCAGAUCCACAC UTT	551
caveolin 2-3	GUUCCUGACGGUGUUCC UGTT	540	CAGGAACACCGUCAGGAA CTT	552
caveolin 2-4	UUGCGGGAAUUCUCUUU GCTT	541	GCAAAGAGAAUUCCCGCA ATT	553
ephA2-1	GGAAGUGGUACUGCUGG ACTT	542	GUCCAGCAGUACCACUUC CTT	554
ephA2-2	CUUCCAGAAGCGCCUGU UCTT	543	GAACAGGCGCUUCUGGAA GTT	555
ephA2-3	GAGCCCCGUAUGCACUG UGTT	544	CACAGUGCAUACGGGGCU CTT	556
ephA2-4	CUACACCUUCACCGUGGA GTT	545	CUCCACGGUGAAGGUGUA GTT	557

Transfection of post-quiescent A549 cells With AntiSense Oligonucleotides.

Materials needed:

- A549 cells can be maintained in DMEM with high glucose (Gibco-BRL) supplemented with 10% Fetal Bovine Serum, 2mM L-Glutamine, and 1X penicillin/streptomycin.
- Opti-MEM (Gibco-BRL)
- Lipofectamine 2000 (Invitrogen)
- Antisense oligomers (Qiagen)
- Polystyrene tubes.
- Tissue culture treated plates.

Quiescent cells are prepared as follows:

Day 0: 300, 000 A549 cells are seeded in a T75 tissue culture flask in 10 ml of A549 media (as specified above), and incubated in at 37°C, 5% CO₂ in a humidified incubator for 48 hours.

Day 2: The T75 flasks are rocked to remove any loosely adherent cells, and the A549 growth media removed and replenished with 10 ml of fresh A549 media. The cells are cultured for six days without changing the media to create a quiescent cell population.

Day 8: Quiescent cells are plated in multi-well format and transfected with antisense oligonucleotides.

A549 cells are transfected according to the following:

1. Trypsinize T75 flask containing quiescent population of A549 cells.
2. Count the cells and seed 24-well plates with 60K quiescent A549 cells per well.
3. Allow the cells to adhere to the tissue culture plate (approximately 4 hours).
4. Transfect the cells with antisense and control oligonucleotides according to the following:

- a. A 10X stock of lipofectamine 2000 (10 ug/ml is 10X) may be prepared, and diluted lipid is allowed to stand at RT for 15 minutes. Stock solution of lipofectamine 2000 is 1 mg/ml. 10 X solution for transfection is 10 ug/ml.
- 5 To prepare 10X solution, dilute 10 ul of lipofectamine 2000 stock per 1 ml of Opti-MEM (serum free media).
- b. A 10X stock of each oligomer may be prepared for use in the transfection. Stock solutions of oligomers are at 100 uM in 20 mM HEPES, pH 7.5. 10X concentration of oligomer may be 0.25 uM.
- 10 To prepare the 10X solutions, dilute 2.5 ul of oligomer per 1 ml of Opti-MEM.
- c. Equal volumes of the 10X lipofectamine 2000 stock and the 10X oligomer solutions are mixed well, and incubated for 15 minutes at RT to allow complexation of the oligomer and lipid. The resulting mixture is 5X.
- 15 d. After the 15 minute complexation, 4 volumes of full growth media is added to the oligomer/lipid complexes (solution may be 1X).
- e. The media may be aspirated from the cells, and 0.5 ml of the 1X oligomer/lipid complexes added to each well.
- 20 f. The cells are incubated for 16-24 hours at 37°C in a humidified CO₂ incubator.
- g. Cell pellets are harvested for RNA isolation and TaqMan analysis of the expression of the protein tyrosine kinase biomarker polypeptides to assess level of knock-down.
- 25

TaqMan Reactions

Quantitative RT-PCR analysis may be performed on total RNA preps that are treated with DNaseI or poly A selected RNA. The Dnase treatment may be performed using methods known in the art, though preferably using a Qiagen RNeasy kit to purify the RNA samples, wherein DNase I treatment is performed on the column.

30

Briefly, a master mix of reagents may be prepared according to the following table:

Dnase I Treatment

<u>Reagent</u>	<u>Per r'xn (in uL)</u>
10x Buffer	2.5
Dnase I (1 unit/ul @ 1 unit per ug sample)	2
DEPC H ₂ O	0.5
RNA sample @ 0.1 ug/ul	20
(2-3 ug total)	
Total	25

5

Next, 5 ul of master mix may be aliquoted per well of a 96-well PCR reaction plate (PE part # N801-0560). RNA samples are adjusted to 0.1 ug/ul with DEPC treated H₂O (if necessary), and 20 ul may be added to the aliquoted master mix for a final reaction volume of 25 ul.

10 The wells are capped using strip well caps (PE part # N801-0935), placed in a plate, and briefly spun in a plate centrifuge (Beckman) to collect all volume in the bottom of the tubes. Generally, a short spin up to 500rpm in a Sorvall RT is sufficient

The plates are incubated at 37°C for 30 mins. Then, an equal volume of 0.1mM EDTA in 10mM Tris may be added to each well, and heat inactivated at 70°C
15 for 5 min. The plates are stored at -80°C upon completion.

RT reaction

A master mix of reagents may be prepared according to the following table:

20

RT Reaction

<u>Reagent</u>	<u>RT</u> <u>Per Rx'n (in ul)</u>	<u>No RT</u> <u>er Rx'n (in ul)</u>
10x RT buffer	5	2.5
MgCl ₂	11	5.5
DNTP mixture	10	5
Random Hexamers	2.5	1.25
Rnase inhibitors	1.25	0.625

<u>Reagent</u>	<u>RT</u> <u>Per Rx'n (in ul)</u>	<u>No RT</u> <u>Per Rx'n (in ul)</u>
RT enzyme	1.25	-
Total RNA 500ng (100ng no RT)	19.0 max	10.125 max
DEPC H ₂ O	-	-
Total	50uL	25uL

Samples are adjusted to a concentration so that 500ng of RNA is added to each RT rx'n (100ng for the no RT). A maximum of 19 ul can be added to the RT rx'n mixture (10.125 ul for the no RT.) Any remaining volume up to the maximum values
 5 may be filled with DEPC treated H₂O, so that the total reaction volume is 50 ul (RT) or 25 ul (no RT).

On a 96-well PCR reaction plate (PE part # N801-0560), 37.5 ul of master mix may be aliquoted (22.5 ul of no RT master mix), and the RNA sample added for a total reaction volume of 50ul (25 ul, no RT). Control samples are loaded into two or
 10 even three different wells in order to have enough template for generation of a standard curve.

The wells are capped using strip well caps (PE part # N801-0935), placed in a plate, and spin briefly in a plate centrifuge (Beckman) to collect all volume in the bottom of the tubes. Generally, a short spin up to 500rpm in a Sorvall RT is
 15 sufficient.

For the RT-PCR reaction, the following thermal profile may be used:

- 25°C for 10 min
- 48°C for 30 min
- 20 • 95°C for 5 min
- 4°C hold (for 1 hour)
- Store plate @-20°C or lower upon completion.

TaqMan reaction (Template comes from RT plate.)

A master mix may be prepared according to the following table:

TaqMan reaction (per well)	
<u>Reagent</u>	<u>Per Rx'n (in ul)</u>
TaqMan Master Mix	4.17
100 uM Probe	.025
100 uM	.05
Forward	
primer	
100 uM	.05
Reverse	
primer	
Template	-
DEPC H ₂ O	18.21
Total	22.5

5

Appropriate forward, reverse, and probe primers may be designed for each protein tyrosine kinase biomarker polypeptides coding region for use in the RT-PCR reaction

10 Using a Gilson P-10 repeat pipetter, 22.5 ul of master mix is aliquoted per well of a 96-well optical plate. Then, using P-10 pipetter, 2.5 ul of sample is added to individual wells. Generally, RT samples are run in triplicate with each primer/probe set used, and no RT samples are run once and only with one primer/probe set, often gapdh (or other internal control).

15 A standard curve is then constructed and loaded onto the plate. The curve has five points plus one no template control (NTC, =DEPC treated H₂O). The curve may be made with a high point of 50 ng of sample (twice the amount of RNA in unknowns), and successive samples of 25, 10, 5, and 1 ng. The curve may be made from a control sample(s) (see above).

20 The wells are capped using optical strip well caps (PE part # N801-0935), placed in a plate, and spun in a centrifuge to collect all volume in the bottom of the tubes. Generally, a short spin up to 500rpm in a Sorvall RT is sufficient.

Plates are loaded onto a PE 5700 sequence detector making sure the plate is aligned properly with the notch in the upper right hand corner. The lid may be tightened down and run using the 5700 and 5700 quantitation program and the SYBR probe using the following thermal profile:

5

- 50°C for 2 min
- 95°C for 10 min
- and the following for 40 cycles:
 - 95°C for 15 sec
 - 60°C for 1 min
- Change the reaction volume to 25ul.

10

Once the reaction may be complete, a manual threshold of around 0.1 may be set to minimize the background signal. Additional information relative to operation of the GeneAmp 5700 machine may be found in reference to the following manuals:

15 “GeneAmp 5700 Sequence Detection System Operator Training CD”; and the “User’s Manual for 5700 Sequence Detection System”; available from Perkin-Elmer and hereby incorporated by reference herein in their entirety.

The skilled artisan would acknowledge that modifications to the above protocol may be required for each protein tyrosine kinase biomarker polypeptide of the present invention. The skilled artisan would also acknowledge that cell lines other than A549 could be used and that A549 are only provided as an example. The skilled artisan would also acknowledge that other means may be used to assess the ability of a complimentary oligonucleotide, such as the RNAi reagents provided in SEQ ID

25 NO:534 to 557, which include, but are not limited to western blots and ELISA assays, among others.

EXAMPLE 6 – ALTERNATIVE METHOD OF ASSESSING ABILITY OF
COMPLIMENTARY SEQUENCES TO MODULATE EXPRESSION LEVELS OF
30 THE PROTEIN TYROSINE KINASE BIOMARKER POLYPEPTIDES OF THE
PRESENT INVENTION.

Preferred complimentary sequences that may be assessed for their ability to modulate the expression levels the protein tyrosine kinase biomarker polypeptides of the present invention are provided as SEQ ID NO:534 to 557. Other complimentary sequences may be designed based upon the coding region of the protein tyrosine kinase biomarker polypeptides of the present invention as provided as SEQ ID NO:1 thru 137, and are specifically contemplated by the present invention.

Co-Transfection RNAi

10 Transfection:

Day prior to transfection, seed 2×10^5 HeLa cells per well of a 24 well dish. The following day, cells should be 90-95% confluent. Dilute 4.5uL of 20uM stock RNAi (one or more of SEQ ID NO:534 to 557) in 50uL Optimem in a polystyrene tube for each RNAi to be transfected (tube A). Mix by gentle tapping. In another polystyrene tube combine 2uL Lipofectamine 2000 with 50 uL Optimem (tube B). Mix by gentle tapping. Allow to sit at RT for 5'. Combine 50uL tube B with the 50uL for each tube A. Mix by gentle tapping. Allow to sit at RT for 15'. Add 500uL serum/antibiotic-free MEM to each tube to give a final RNAi concentration of 150nM. (For cotransfections of RNAi with plasmid, use 1uL of 20uM stock RNAi (final concentration of 33nM) along with 1ug vector DNA in tube A, and then proceed with transfection protocol above). Remove the media from HeLa plates and replace with the 600uL transfection mix. Put in 37°C 5% CO2 incubator for 4-5 hours. Replace the media with MEM containing 10% FBS.

25 Controls to include in the transfection include a fluorescent oligonucleotide control (1U/uL=20uM) to calculate transfection efficiency, GFP B as a non-specific negative control, CDC2 as a normalizing knockdown control, and an untransfected control receiving no DNA.

30 Lysis:

48 hours post-transfection, aspirate media and wash cells 1X with approx. 500uL cold 1xPBS per well. Aspirate and replace with 100uL cold RIPA containing protease inhibitors (1 mini BM protease inhibitor tablet/10mL 1x RIPA). Rock and tap the plate a few times and place at 4°C for 10-15 minutes. Tap/rock the plate several more times. Using a 200uL pipetteman, aspirate 5-10 times and wash the well to ensure complete lysis and transfer of all material. Transfer lysate to an eppendorf tube and pipette up and down 5-10 times. If sample is still viscous, pipette up and down several more times. Spin samples down for 10' at 14000 RPM 4°C. Samples can now be stored at -20°C or prepared for loading.

10

Western blotting/Novex:

Prepare sample by combining 20uL lysate with 3uL reducing reagent and 7uL 4X gel loading dye. Heat at 70°C for 10' and then place samples on ice. While samples are heating, prepare desired gel (usually a 4-12% Bis-Tris gel) by removing comb and sealing tape. Place gels in gel box and fill inner and outer chambers with desired buffer (either 1x MES or MOPS- Add 50mL 20x buffer to 950mL dH2O for each gel box).. Add 600uL Oxidizing reagent to the inner chamber. Wash out each well by blasting with 500uL buffer. In well one, load 5uL marker- Invitrogen's SeeBlue Plus2. Load samples in subsequent lanes. Run gel at 200V for 45-50 minutes. Make up 1X transfer buffer- 50mL 20x transfer buffer, Methanol (100mL if transferring one gel, 200mL if transferring 2 gels in the same apparatus) and dH2O to 1000mL. Soak blotting pads in dH2O and then transfer buffer- make sure to push down on pads to rid of air bubbles. Soak precut Hybond-ECL membrane (Amersham nitrocellulose) in dH2O and then in transfer buffer. Cut the end off of Biorad filter paper to match size of transfer membrane. If transferring one gel, place 2 blotting pads into blotting chamber. For 2 gels, place down 1 pad. Briefly soak a filter paper in transfer buffer and carefully lay on blotting pad. Open gel cassette with cracking tool, cut off top, bottom and sides of gel. Briefly rinse it in transfer buffer and then lay it on filter paper carefully making sure no air bubbles are present. Lay transfer membrane on top again being careful there are no bubbles. Put down filter paper. Put down 2 blotting pads if transferring one gel to complete the sandwich. If transferring

25

30

2, put down 1 blotting pad, filter paper, gel, membrane, filter paper, blotting pad.

Gels are now ready for transfer. Squeeze together the gel sandwich and place in transfer apparatus. Fill inner and outer chambers with transfer buffers. Transfer gels for 1 hour at 30V.

- 5 Remove membranes and place them in Superblock (Pierce) and rock at RT for a minimum of 1 hour-overnight. (I have stored membranes in Superblock at 4°C over the weekend). Primary antibody and normalizing antibody are diluted in a 1:10 mix of Superblock:1xPBS/0.3% Tween-20. Membranes are incubated and rocked at RT in primary antibody for a minimum of 1 hour-overnight. Membranes are then
- 10 washed thoroughly in 1xPBS/0.3% Tween-20. I usually give several quick rinses and then rinse 3X 5' in 1xPBS/0.3% Tween-20. During the final wash, HRP-conjugated secondary antibody is diluted in 1xPBS/0.3% Tween-20. Add this to membrane and rock at RT for a minimum of 30'. Wash membranes thoroughly. I usually give several quick rinses and then rinse 3X 5' in 1xPBS/0.3% Tween-20. Membranes are
- 15 removed from wash buffer and the excess buffer drained by holding the edge of the membranes on a paper towel. Membranes are placed on Saran Wrap that has been smoothed on benchtop to remove air bubbles. Enough ECL reagent is added to cover the membrane for 1 minute. Remove membranes and drain off excess ECL on paper towel. Place membranes in between two transparency sheets, being careful to smooth
- 20 out air bubbles.

Quantitation:

- Expose membranes using FluorS-Max. Relative percent inhibition may be
- 25 determined by comparing the intensity of each band with RNAi treatment to the intensity of each band without RNAi treatment (control). Normalize lanes by dividing band of interest by normalizing band for each lane. Divide the normalized value for each treated sample by the normalized value of the control. Percent inhibition can then be calculated by using the formula $(1 - \text{above value}) \times 100$.

- 30 The skilled artisan would acknowledge that modifications to the above protocol may be required for each protein tyrosine kinase biomarker polypeptide of the present invention.

2, put down 1 blotting pad, filter paper, gel, membrane, filter paper, blotting pad. Gels are now ready for transfer. Squeeze together the gel sandwich and place in transfer apparatus. Fill inner and outer chambers with transfer buffers. Transfer gels for 1 hour at 30V.

- 5 Remove membranes and place them in Superblock (Pierce) and rock at RT for a minimum of 1 hour-overnight. (I have stored membranes in Superblock at 4⁰C over the weekend). Primary antibody and normalizing antibody are diluted in a 1:10 mix of Superblock:1xPBS/0.3% Tween-20. Membranes are incubated and rocked at RT in primary antibody for a minimum of 1 hour-overnight. Membranes are then
- 10 washed thoroughly in 1xPBS/0.3% Tween-20. I usually give several quick rinses and then rinse 3X 5' in 1xPBS/0.3% Tween-20. During the final wash, HRP-conjugated secondary antibody is diluted in 1xPBS/0.3% Tween-20. Add this to membrane and rock at RT for a minimum of 30'. Wash membranes thoroughly. I usually give several quick rinses and then rinse 3X 5' in 1xPBS/0.3% Tween-20. Membranes are
- 15 removed from wash buffer and the excess buffer drained by holding the edge of the membranes on a paper towel. Membranes are placed on Saran Wrap that has been smoothed on benchtop to remove air bubbles. Enough ECL reagent is added to cover the membrane for 1 minute. Remove membranes and drain off excess ECL on paper towel. Place membranes in between two transparency sheets, being careful to smooth
- 20 out air bubbles.

Quantitation:

- Expose membranes using FluorS-Max. Relative percent inhibition may be
- 25 determined by comparing the intensity of each band with RNAi treatment to the intensity of each band without RNAi treatment (control). Normalize lanes by dividing band of interest by normalizing band for each lane. Divide the normalized value for each treated sample by the normalized value of the control. Percent inhibition can then be calculated by using the formula $(1 - \text{above value}) \times 100$.

- 30 The skilled artisan would acknowledge that modifications to the above protocol may be required for each protein tyrosine kinase biomarker polypeptide of the present invention.

BRIEF DESCRIPTION OF THE SEQUENCE LISTING

Incorporated herein by reference in its entirety is a Sequence Listing, comprising SEQ ID NO:1 through SEQ ID NO:557, which include nucleic acid and amino acid sequences of the protein tyrosine kinase biomarkers as presented in Table 2 herein and the nucleotide sequences of forward and reverse primer pairs for the polynucleotide markers, probes, and RNAi reagents as described herein. The Sequence Listing is contained on a compact disc, i.e., CD-ROM, three identical copies of which are filed herewith. The Sequence Listing, in IBM/PC MS-DOS text format, was first created on August 25, 2003, and is 896 KB in size.

10

The contents of all patents, patent applications, published PCT applications and articles, books, references, reference manuals, abstracts, the Sequence Listing, and internet websites cited herein are hereby incorporated by reference in their entirety to more fully describe the state of the art to which the invention pertains.

15

As various changes can be made in the above-described subject matter without departing from the scope and spirit of the present invention, it is intended that all subject matter contained in the above description, or defined in the appended claims, be interpreted as descriptive and illustrative of the present invention. Many modifications and variations of the present invention are possible in light of the above teachings.

20

WHAT IS CLAIMED IS:

1. A predictor set comprising a plurality of polynucleotides whose expression pattern is predictive of the response of cells to treatment with a compound that modulates protein tyrosine kinase activity or members of the protein tyrosine kinase pathway.
5
2. The predictor set according to claim 1 wherein the polynucleotides are selected from the group consisting of:
 - a.) the polynucleotides provided in Table 2;
 - b.) the sensitive predictor polynucleotides provided in
10 Table 2; and
 - c.) the resistant predictor polynucleotides provided in Table 2.
3. The predictor set according to claim 2 wherein the plurality of polynucleotides comprise the number of polynucleotides selected from the group consisting of:
15
 - a.) at least about 1 polynucleotides;
 - b.) at least about 3 polynucleotides;
 - c.) at least about 5 polynucleotides;
 - d.) at least about 7 polynucleotides;
 - e.) at least about 10 polynucleotides;
 - f.) at least about 15 polynucleotides;
 - 20 g.) at least about 20 polynucleotides;
 - h.) at least about 25 polynucleotides; and
 - i.) at least about 30 polynucleotides.
4. The predictor set according to claim 3 wherein the plurality of polynucleotides comprise a member of the group consisting of:
25
 - a.) the polynucleotides provided in Table 3;
 - b.) the sensitive predictor polynucleotides provided in
30 Table 3;

- 5
- c.) the resistant predictor polynucleotides provided in Table 3;
 - d.) the polynucleotides provided in Table 4;
 - e.) the sensitive predictor polynucleotides provided in Table 4;
 - f.) the resistant predictor polynucleotides provided in Table 4;
 - g.) the polynucleotides provided in Table 5;
 - h.) the sensitive predictor polynucleotides provided in Table 5; and
 - 10 i.) the resistant predictor polynucleotides provided in Table 5.

15 5. The predictor set according to claim 4 wherein the compound is selected from the group consisting of:

- a.) antisense reagents directed to said polynucleotides;
- b.) antibodies directed against polypeptides encoded by said polynucleotides; and
- c.) small molecule compounds.

20 6. The predictor set according to claim 5 wherein the compound is BMS-A.

7. The predictor set according to claim 1 wherein said cells are a member of the group consisting of: breast cells, and breast cancer cells.

25 8. A predictor set comprising a plurality of polypeptides whose expression pattern is predictive of the response of cells to treatment with compounds that modulate protein tyrosine kinase activity or members of the protein tyrosine kinase pathway.

9. The predictor set according to claim 8 wherein the polypeptides are selected from the group consisting of:

- 30
- a.) the polypeptides provided in Table 2;
 - b.) the sensitive predictor polypeptides provided in Table 2; and

- c.) the resistant predictor polypeptides provided in Table 2.

10. The predictor set according to claim 9 wherein the plurality of polypeptides comprise the number of polypeptides selected from the group consisting of:

5

- a.) at least about 1 polypeptides;
- b.) at least about 3 polypeptides;
- c.) at least about 5 polypeptides;
- d.) at least about 7 polypeptides;
- 10 e.) at least about 10 polypeptides;
- f.) at least about 15 polypeptides;
- g.) at least about 20 polypeptides;
- h.) at least about 25 polypeptides; and
- i.) at least about 30 polypeptides.

15

11. The predictor set according to claims 10 wherein the plurality of polypeptides comprise a member of the group consisting of:

20

- a.) polypeptides provided in Table 3;
- b.) the sensitive predictor polypeptides provided in Table 3;
- c.) the resistant predictor polypeptides provided in Table 3;
- d.) the polypeptides provided in Table 4;
- e.) the sensitive predictor polypeptides provided in Table 4;
- f.) the resistant predictor polypeptides provided in Table 4;
- g.) the polypeptides provided in Table 5;
- h.) the sensitive predictor polypeptides provided in Table 5;
- 25 and
- i.) the resistant predictor polypeptides provided in Table 5.

12. The predictor set according to claim 11 wherein the compound is selected from the group consisting of:

30

- a.) antisense reagents directed against polynucleotides encoding said polypeptides;
- b.) antibodies directed against said polypeptides; and

c.) small molecule compounds.

13. The predictor set according to claim 12 wherein the compound is BMS-A.

5 14. The predictor set according to claim 8 wherein said cells are a member of the group consisting of: breast cells, and breast cancer cells.

15. A method for predicting whether a compound is capable of modulating the activity of cells, comprising the steps of:

- 10 a.) obtaining a sample of cells;
b.) determining whether said cells express a plurality of markers; and
c.) correlating the expression of said markers to the compounds ability to modulate the activity of said cells.

15 16. The method according to claim 15 wherein the plurality of markers are polynucleotides.

17. The method according to claim 16 wherein the polynucleotides are the polynucleotides of claim 4.

20 18. The method according to claim 17 wherein the compounds are a member of the group consisting of:

- a.) the compounds according to claim 5; and
b.) the compounds according to claim 6.

25 19. The method according to claim 18 wherein the cells are a member of the group consisting of: breast cells, and breast cancer cells.

20. The method according to claim 15 wherein the plurality of markers are polypeptides.

21. The method according to claim 20 wherein the polypeptides are the polypeptides of claim 11.

30 22. The method according to claim 21 wherein the compounds are a member of the group consisting of:

- c.) the compounds according to claim 12; and

d.) the compounds according to claim 13.

23. The method according to claim 19 wherein the cells are a member of the group consisting of: breast cells, and breast cancer cells.
- 5 24. A plurality of cell lines for identifying polynucleotides and polypeptides whose expression levels correlate with compound sensitivity or resistance of cells associated with a disease state.
- 10 25. The plurality of cell lines according to claim 24 wherein said cell lines are breast cancer cell lines.
26. The plurality of cell lines according to claim 25 wherein said cell lines comprise one or more cell lines provided in Table 1.
- 15 27. A method of identifying polynucleotides and polypeptides that predict compound sensitivity or resistance of cells associated with a disease state, comprising the steps of:
- a.) subjecting the plurality of cell lines according to claim 26 to one or more compounds;
 - 20 b.) analyzing the expression pattern of a microarray of polynucleotides or polypeptides; and
 - c.) selecting polynucleotides or polypeptides that predict the sensitivity or resistance of cells associated with a disease state by using said expression pattern of said microarray.
- 25 28. The method according to claim 24 wherein the compounds are a member of the group consisting of:
- a.) the compounds according to claim 5; and
 - b.) the compounds according to claim 6;
 - c.) the compounds according to claim 12; and
 - 30 d.) the compounds according to claim 13.

29. The method according to claim 29 wherein said disease is breast cancer.

30. A method for predicting whether an individual requiring treatment for
5 a disease state, will successfully respond or will not respond to said treatment comprising the steps of:

- a.) obtaining a sample of cells from said individual;
- b.) determining whether said cells express a plurality of markers; and
- 10 c.) correlating the expression of said markers to the individuals ability to respond to said treatment.

31. The method according to claim 30 wherein the plurality of markers are polynucleotides.

15 32. The method according to claim 31 wherein the polynucleotides are the polynucleotides of claim 4.

33. The method according to claim 32 wherein the compounds are a member of the group consisting of:

- a.) the compounds according to claim 5; and
- 20 b.) the compounds according to claim 6.

34. The method according to claim 33 wherein the disease state is breast cancer.

25 35. The method according to claim 30 wherein the plurality of markers are polypeptides.

36. The method according to claim 35 wherein the polypeptides are the polypeptides of claim 11.

37. The method according to claim 36 wherein the compounds are a member of the group consisting of:

- 30 a.) the compounds according to claim 5; and
- b.) the compounds according to claim 6.

38. The method according to claim 37 wherein the disease state is breast cancer.
39. A method of screening for candidate compounds capable of binding to and/or modulating the activity of a protein tyrosine kinase biomarker polypeptide, comprising:
- 5 (a) contacting a test compound with a polypeptide according to claim 11; and
- (b) selecting as candidate compounds those test compounds that bind to and/or modulate activity of the polypeptide.
- 10 40. A method of treating breast cancer in a subject, comprising administering a modulator of one or more protein tyrosine kinase biomarker polypeptides, wherein said polypeptide(s) is selected from the group consisting of:
- a.) polypeptides provided in Table 2;
- b.) the sensitive predictor polypeptides provided in Table 2;
- 15 c.) the resistant predictor polypeptides provided in Table 2;
- d.) polypeptides provided in Table 3;
- e.) the sensitive predictor polypeptides provided in Table 3;
- f.) the resistant predictor polypeptides provided in Table 3;
- g.) the polypeptides provided in Table 4;
- 20 h.) the sensitive predictor polypeptides provided in Table 4;
- i.) the resistant predictor polypeptides provided in Table 4;
- j.) the polypeptides provided in Table 5; and
- k.) the sensitive predictor polypeptides provided in Table 5.

FIG. 1

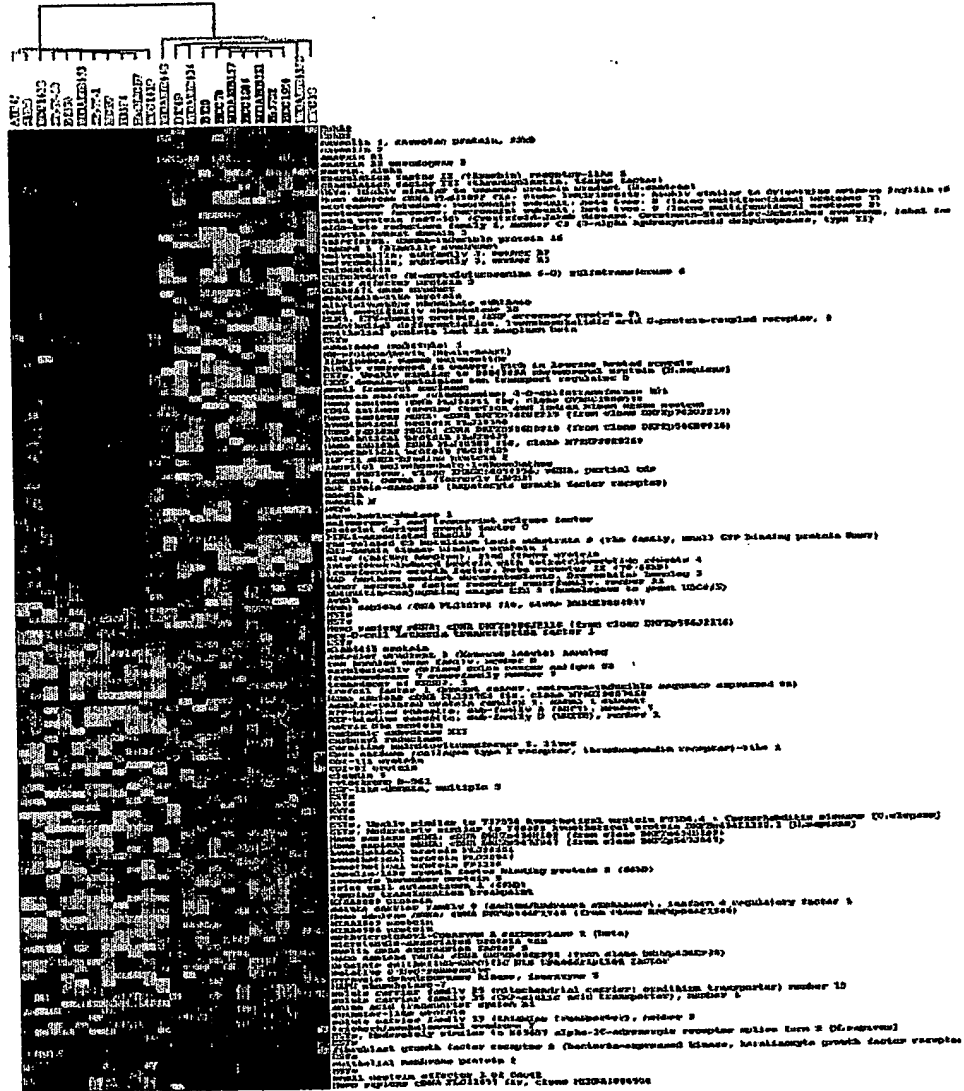
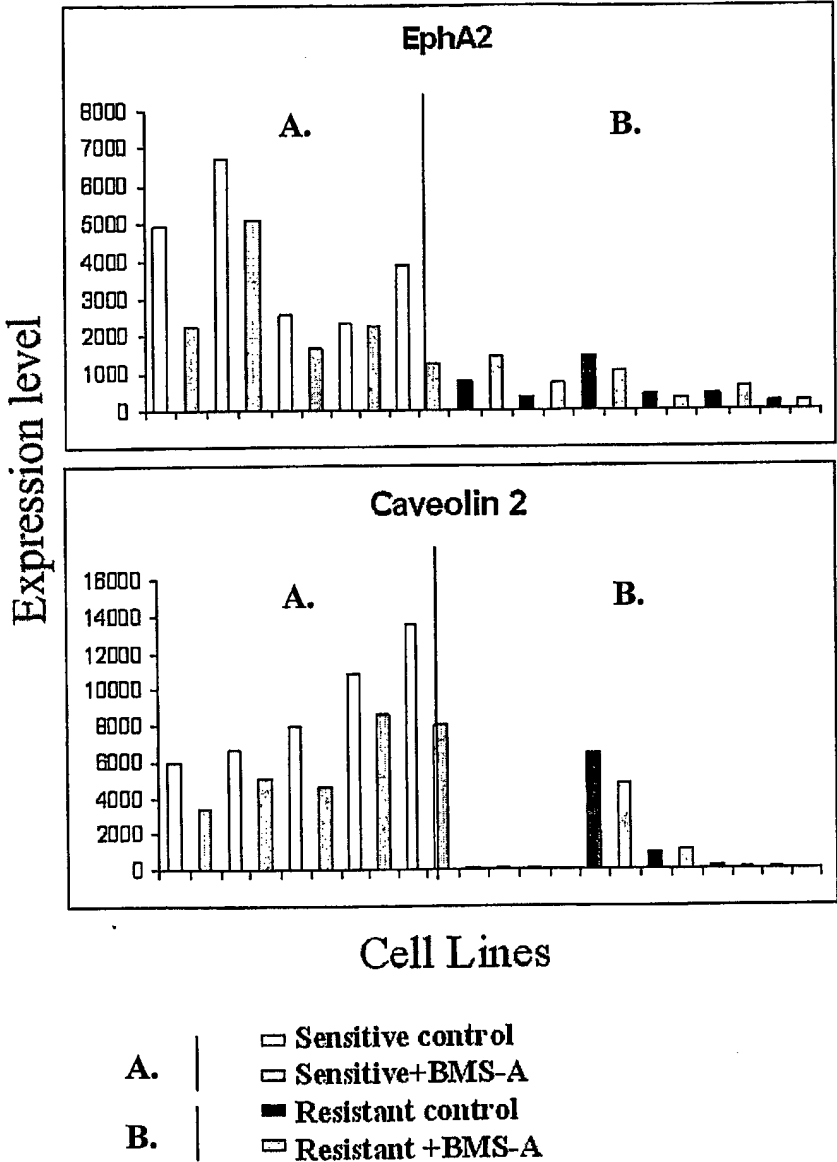
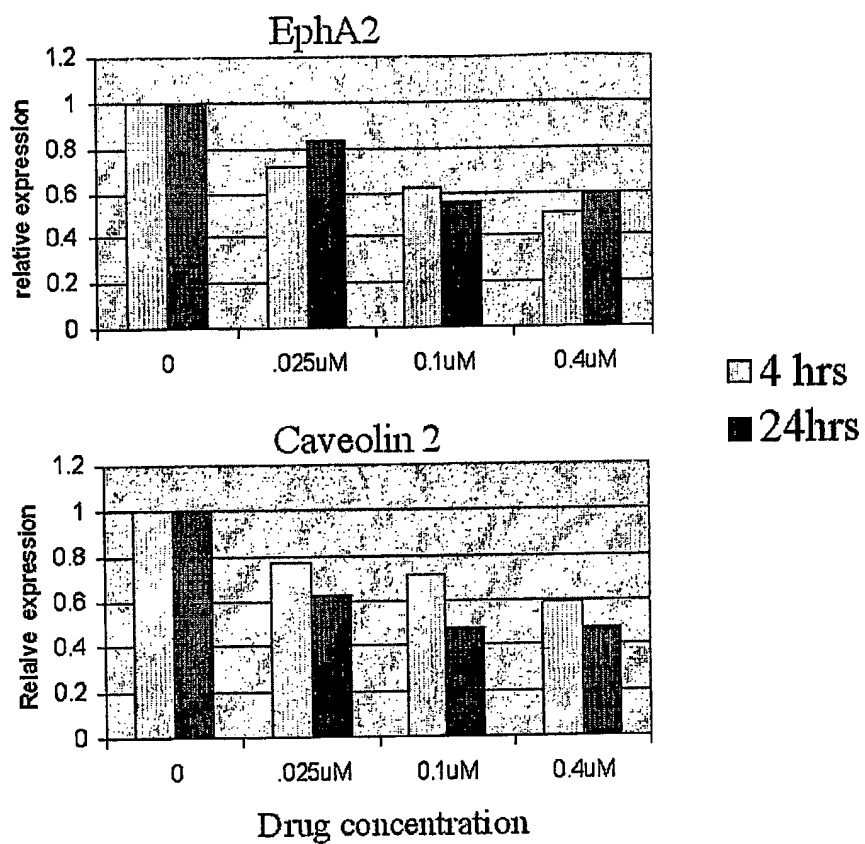


FIG. 2



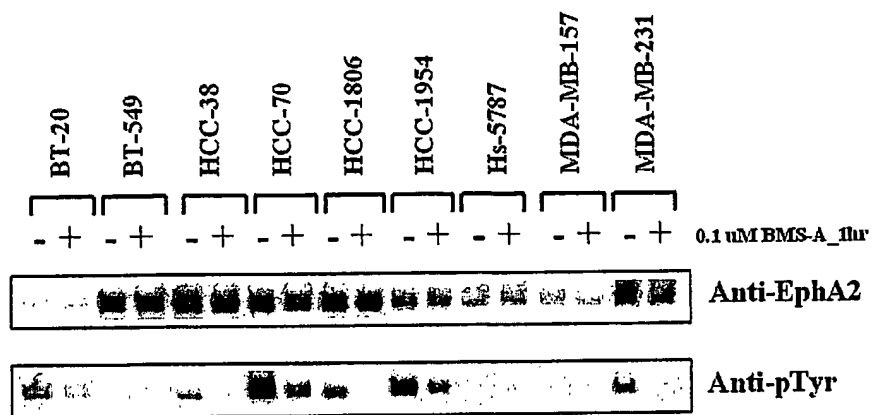
3/7

FIG. 3



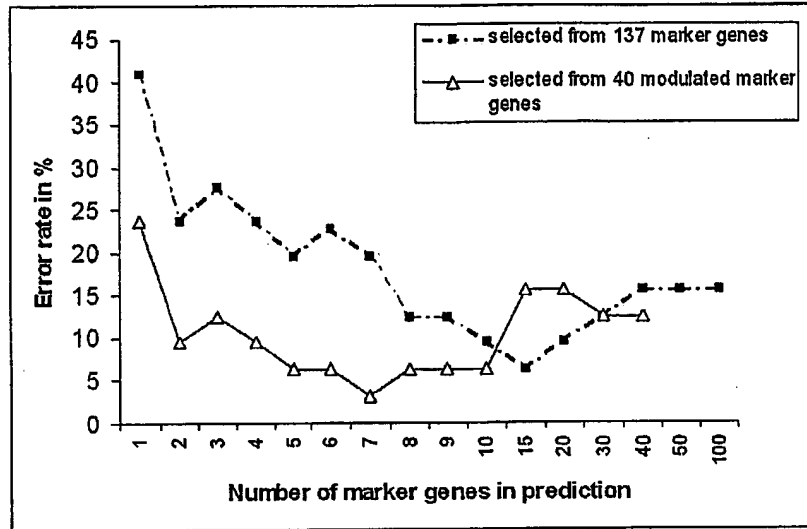
4/7

FIG. 4



5/7

FIG. 5



6/7

FIG. 6

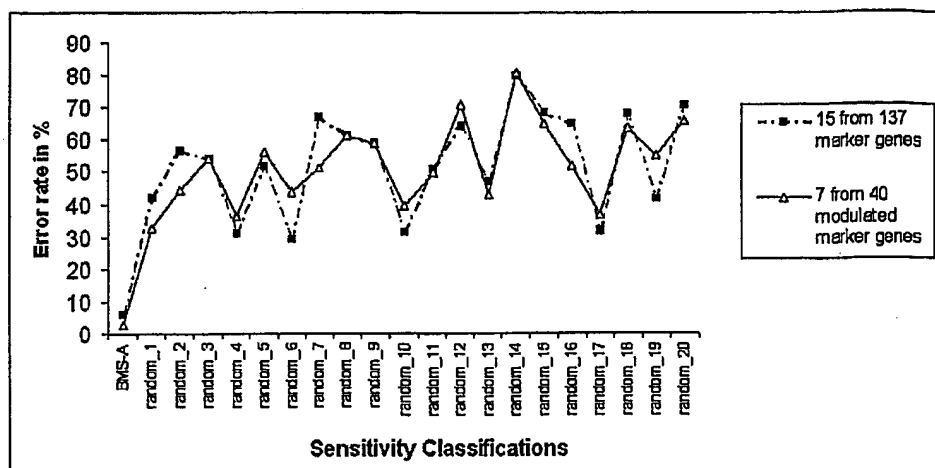
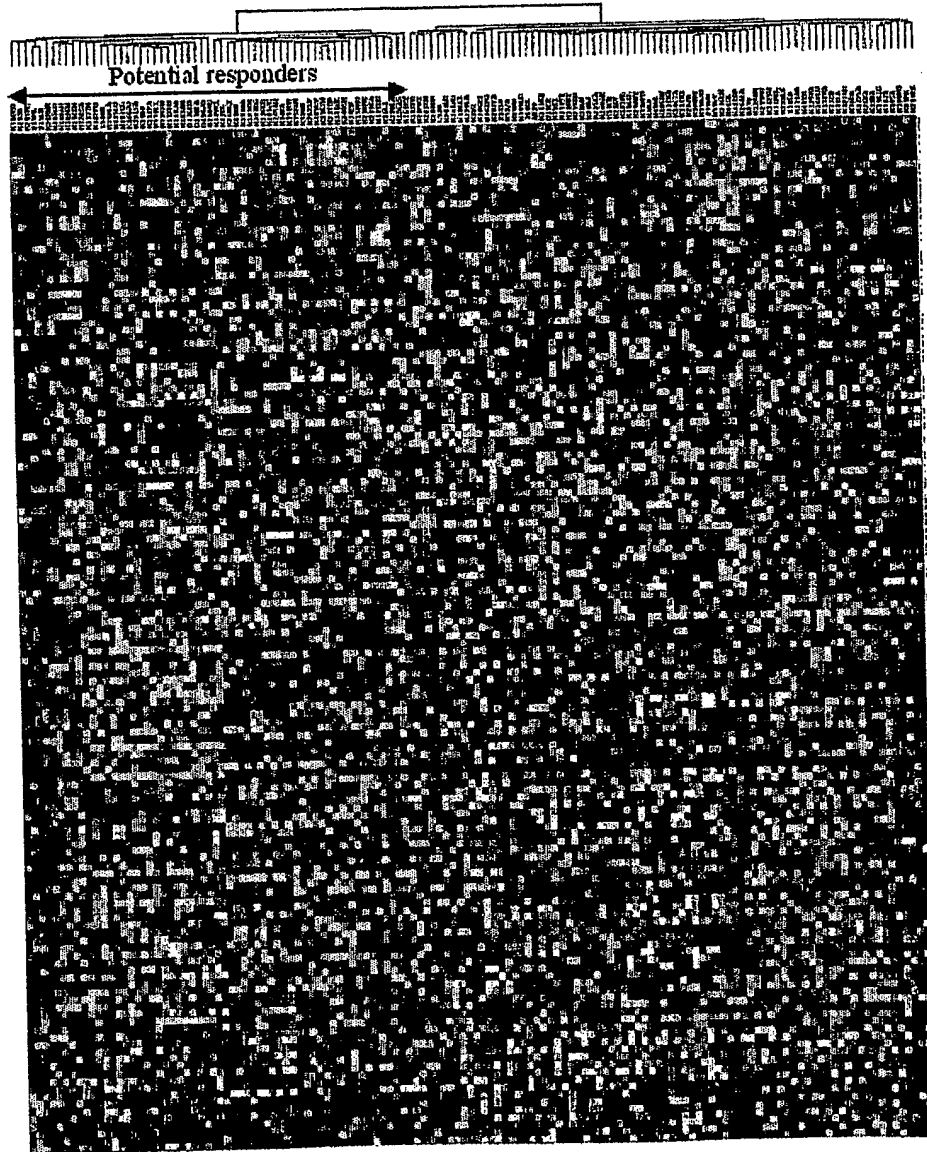


FIG. 7



SEQUENCE LISTING

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INTERACT WITH AND/OR MODULATE PROTEIN TYROSINE KINASES AND/OR
PROTEIN TYROSINE KINASE PATHWAYS IN BREAST CELLS

<130> D0273 PCT

<150> 60/406,385

<151> 2002-08-27

<160> 557

<170> PatentIn version 3.2

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<212> DNA

<213> Homo sapiens

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1042

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 <212> DNA
 <213> Homo sapiens

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<210> 68
<211> 2201
<212> DNA
<213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

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<210> 70
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<212> DNA
<213> Homo sapiens

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cctcagttcc tgtgcctcag cccacaggcc actgtgataa tgggtctgtt agcacttctg 360
tattttattgt aagaatgctt atcatgaaga tacacactgt aactacaaga aattataaat 420
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<210> 71

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<211> 456
 <212> DNA
 <213> Homo sapiens

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<210> 72
 <211> 2278
 <212> DNA
 <213> Homo sapiens

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<212> DNA
<213> Homo sapiens

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1819

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<211> 871

<212> DNA

<213> Homo sapiens

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 <211> 956
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<210> 77
 <211> 591
 <212> DNA
 <213> Homo sapiens

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 <212> DNA
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 <212> DNA
 <213> Homo sapiens

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<212> DNA
<213> Homo sapiens

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atgta

605

<210> 101

<211> 950

<212> DNA

<213> Homo sapiens

<400> 101

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tcaccaccaa gacgcagccc gtggaagcca ccgatgatgc cttttgggac cagttctggg      180
cagacacagc cacctcgggtg caggatgtgt ttgcaactgt gccggcagca gagatccggg      240
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tgggtgcaggg agctgagagt ggctgccact cggagaagga gaagcagatc gtcctgaact      360
gcagccgggt gctcaccgcg gtgctgccct acatctttga ggaccccgac tggaggggct      420
tcttctggtc cacagtgtcc ggggcagggc gaggagggca gggagaagag gatgatgagc      480
atgccaggcc cctggccgag tccctgctcc tggccattgc tgacctgtc ttctgcccgg      540
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ggacagctgt gaatacatct gggaggctgg tgtgggcttc gtcactccc cccagcctaa      660
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<210> 102

<211> 961

<212> DNA

<213> Homo sapiens

<400> 102

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aaggaatgca ttaccacatg accacatgct gagaccccat ggggtctaac acgggaccta      180
agaaagtctc tgcagccaga tagtacatgg tgtctccaca aaactaggca ttctggagat      240
tgcccagaaa gggatgtgag gggaccgtta agatctgtct tgcttatctc atgcactcac      300

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attccttcag cctcctggag ttcttgataa aaggaagcca ggtgtggaca ttttttagct 360
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ccgtttttct ggaaaaagta gtatgcccat gtatgtgtgt ttttcttaac acaggtccat 480
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<210> 103
<211> 405
<212> DNA
<213> Homo sapiens

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<400> 103
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cagcccgacc agcagaggag gtggcagcac cagggcccca acctgggtgt ccggttctgc 180
tctgggggtca ggctggaggc tggccctctg ggtgcctctg ggaatgaggg ggctgggcta 240
cataactatg gtcccctccc aactctaaca tactagatac tacaacggct ggggaaactc 300
ccgaacagga tcttaacaga gagagggaga gagagaaaga gaggagagag agagagagag 360
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<210> 104
<211> 2051
<212> DNA
<213> Homo sapiens

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<400> 104
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ttgtggatga gttttgtgag agaacagaga gaccattgta cctggcaciaa gggtctttca 180
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 tcagcacagc cctggattgc agccccgagg ctgagaccag acaaagcccg ggaggcagaa 300
 agatgctcca agaaccaaca ctatcaatgt ctttgcaaat cctcacagga ttctgtggg 360
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 gccctgaag gatgccagta ctgtggtgtg tgagtctcag cagccgcca cacgtccta 480
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aaaaaaaaa a 2051

<210> 105
<211> 1291
<212> DNA
<213> Homo sapiens

<400> 105
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gggtttggca tatttgacat ggagccccat gccaaaggat gcagctgttg gtgtctgtcc 180
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1291

<210> 106

<211> 641

<212> DNA

<213> Homo sapiens

<400> 106

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gccacacag tgccctgtctg agtccttctg tgcccaaagtg tggagcaggc tgcagagact	120
tgaagcctgg gggtttgtgc ctcccttttg ttttgttttt tttgagacag agtcttgctc	180
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gttcaagtga ttctcctgcc tcagcctcca gagtagctgg gattacaggt gtgcaccacc	300
acgccaggct aatttttgta ttttagtag agacagggtt tcaccatggt ggccaggctg	360
gtctccaact cctggcctca agtcctcctc ccgccttagc ctcccaaagt gctggaatta	420
caggcatgag ccaccacgcc cgggcacctt cttgccagca actcttagac caggaagcct	480
cggatggcag ctatgaagtc ctgtgggcgt tcagcgtgga tccagtggcc agcgttcagc	540
accgtctgca tctggacagg atggaagagc cgcataatct tcagggtgata gctggaatgc	600
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<210> 107

<211> 1433

<212> DNA

<213> Homo sapiens

<400> 107

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 gaggaagggg gttgtggtcg gggagctggg gtacaggttt ggggaggggg aagagaaatt 1380
 tttatttttg aaccctgtg tcccttttgc ataagattaa aggaaggaaa agt 1433

<210> 108

<211> 1825

<212> DNA

<213> Homo sapiens

<400> 108

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 accagatctg cgcacggggg acggacgtgc ccgggcagat gggggcctac ggggtgacac 180
 cgaggccggg acagcttcag gggccccaga aggacctgac ccagaaattg aggtccccgc 240
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 gcttcgggtcg ttccgatggc agtgagagacc acggtccaca ctcacctctc tgcgtctcca 420
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 tacgcggcgc ccctgggatc gtacccttac ggggaccag cgtaccggaa gaacgccaca 540
 agggacgcca cggctaccct caaggcctgg ctcaacgagc accgcaagaa cccctacccc 600

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accaagggcg agaagatcat gctggccatc atcaccaaga tgaccctcac ccagggtgtcc 660
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gacatttaac gcgggctgcg tcgggtcccg acttttctaa tttattaaaa acatggcctt 1680
ggcagttatt tttccatcac cgagagagag agacagagag agaaaataaa ctaccctcc 1740
tattcagaag tttatagttt atggagatgg atgacataaa aatgtaaaca tctccacaca 1800
cacaaaaaaa tgttttaacc aaccg 1825

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<210> 109
<211> 1427
<212> DNA
<213> Homo sapiens

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<400> 109
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tttctgccc cttctcccca aatcatcaac aatagaagaa gaagaaaaca tgtcaggaca 180
caaatgcagt tatccctggg acttacagga tcgatatgct caagataagt cagttgtaaa 240

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taagatgcaa cagaaatatt gggagacgaa gcaggccttt attaaagcca caggggaagaa 300
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 aatgaaaggg tcaacggccc tgtttatatt ggtataaaaa aaaaaa 1427

<210> 110
 <211> 823
 <212> DNA
 <213> Homo sapiens

<400> 110
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 gactaggcct caggcagaga ctaaaggaca tctcttgggg tgcctgaag tgatttggac 180
 ccctgagggc agacacctaa gtaggaatcc cagtgggaag caaagccata aggaagccca 240
 ggattccttg tgatcaggaa gtgggccagg aaggctctga ccagctcaca tctcaactgc 300

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atgcagcacg gaccggatgc gcccaactggg tcttggttc cctcccatct tctcaagcag   360
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cacttgggtc tccagatttt ctgttacgtc cttgtgggtc aggatatttc tggaagtcac   480
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atgttggtgc tctccagctc ctggactttt ttctgtaatt cttggttctg ttcagaacag   720
gctgccaccc tgctctccag cccatcaatg tactccttct tccgcgcgcg actgtmtga   780
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```

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<210> 111
<211> 553
<212> DNA
<213> Homo sapiens

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<220>
<221> misc_feature
<222> (498)..(498)
<223> n is a, c, g, or t

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<400> 111
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gttccctttg cacaccggca cctcatcaca acaccctctt ggtgtggatg ccatggggcc   180
atgctgtagt caaaagttaa atgaaaaacc acaagtttag tttgactcgc tctcctaggg   240
tggaatttcac tcagatattt gttccatatt ataggagggt ggatcctagc aaggcaacag   300
tgtagttttt acattcacag attggctgaa gtagtacaaa ttgagctgct aatctaggtg   360
tctccctccc tgttaccata cttcataaga aatgtgaatt aaaatgaaca atggaccaca   420
ggtgggttata aaaatagata actcgagag tccataaata tctacagtta gtagagccag   480
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gaggacacag tca                                           553

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<210> 112
<211> 425
<212> DNA
<213> Homo sapiens

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<220>
 <221> misc_feature
 <222> (388)..(388)
 <223> n is a, c, g, or t

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<212> DNA

<213> Homo sapiens

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<213> Homo sapiens

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 <213> Homo sapiens

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<212> DNA
<213> Homo sapiens

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 <213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

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 <211> 487
 <212> DNA
 <213> Homo sapiens

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 gagttatcca actttttttg actcaaagat agtttgotta gatttttttt ttttgagatg 180


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<211> 4574
<212> DNA
<213> Homo sapiens

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cataggggtc ttcttaatcg cctgtatggt ggtaacagtc atcctgtgcc gaatgaagaa	1800
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 aaagtattgt gttttgcttt ggaaacaccc actcactttg caatagccgt gcaagatgaa 4440
 tgcagattac actgatctta tgtgttacaa aattggagaa agtatttaaat aaaacctgtt 4500
 aatttttata ctgacaataa aaatgtttct acagatatta atgttaacaa gacaaaataa 4560
 atgtcacgca actt 4574

<210> 133
 <211> 549
 <212> DNA
 <213> Homo sapiens

<400> 133
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 ccttctatta tttatttact tcttagacac ttgtcgaaat ctgaaaattc tttattcggt 180
 ggtttgtcta tattgacaga ctgtaactgg atgaaggcaa gagaccttgt cttgtacata 240
 gttgtgtcct cagcacttgg cagaatgtct agatggagca ggtgcttaaa tacttgttga 300
 aaaaataaaa actgcaactg caggggtttt taaatatata aacataaaaa tattactcat 360
 caaaatttaa gtaatatagc taaagctatg ctcagaggaa cattcaatag gattacattt 420

ttacattata aaacataaaa tattataaaa gttgcaattt tattaagtta aaaaattcaa 480
 aactggccag gcacagtggg tcacgcctgt aatoccagca ctttgggagg ccgagggtggg 540
 tagatcacc 549

<210> 134
 <211> 799
 <212> DNA
 <213> Homo sapiens

<400> 134
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 acatcccgt ctgggcttta aacgtgaccc ctgcctcga ctgcacctgc cctgtgaaaa 120
 tgttggtgct tcttgctttc atcatcgctt tccacatcac ctctgcagcc ttgctgttca 180
 ttgccaccgt cgacaatgcc tgggtgggtag gagatgagtt ttttgcagat gtctggagaa 240
 tatgtaccaa caacacgaat tgcacagtca tcaatgacag ctttcaagag tactccacgc 300
 tgcaggcggt ccaggccacc atgatectct ccaccattct ctgctgcac gccttcttca 360
 tcttcgtgct ccagctcttc cgctgaagc agggagagag gtttgccta acctccatca 420
 tccagcta atgtcatgtctg tgtgtcatga ttgcggcctc catttataca gacaggcggtg 480
 aagacattca cgacaaaaac gcgaaattct atcccgtgac cagagaaggc agctacggct 540
 actcctacat cctggcggtg gtggccttcg cctgcacctt catcagcggc atgatgtacc 600
 tgatactgag gaagcgcaaa tagagttccg gagctgggtt gcttctgctg cagtacagaa 660
 tccacattca gataaccatt ttgtatataa tcattatttt ttgaggtttt tctagcaaac 720
 gtattgtttc ctttaaaagc caaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaagaa 780
 aaaaaaaaaa aaaaaaaaaa 799

<210> 135
 <211> 561
 <212> DNA
 <213> Homo sapiens

<220>
 <221> misc_feature
 <222> (368)..(368)
 <223> n is a, c, g, or t

<400> 135
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 attatcctat caagggcaga aatgttagat cttactccaa gataggtaaa cgcctttgaa 120

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acgcaacaaa aagagacgat gatcttatga gctcatttat gttcatgcgt gaaagtgtga    180
aggtcactag ctttgctgtg tttctacaag tttccttgac tgtaaaaaca gtcaaaatgt    240
aaccaaccta attcaagatg ttaaattaat taagttaaat aaaattagcc aaagcactgt    300
aaattaaatg acataatact taagttctgt actgatgaca ccctttgatc aaaagaaggt    360
ggacccanta aggtgcttct ggagggttatt acttctctaa ttccgaattt atcatcactg    420
actctttagg cataataaaa tgtagagtgt aaatagggtg gtgacaattt atttaattta    480
ggttgagcta cttataaata cctgagagct ctggaaacaa catatatatc catcttgcaa    540
atatcagtaa aaagaacaag t                                              561

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<210> 136
<211> 567
<212> DNA
<213> Homo sapiens

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<220>
<221> misc_feature
<222> (61)..(61)
<223> n is a, c, g, or t

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<220>
<221> misc_feature
<222> (432)..(432)
<223> n is a, c, g, or t

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<220>
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<223> n is a, c, g, or t

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<220>
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<222> (465)..(465)
<223> n is a, c, g, or t

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<220>
<221> misc_feature
<222> (495)..(495)
<223> n is a, c, g, or t

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<220>
<221> misc_feature
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<223> n is a, c, g, or t

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<220>
<221> misc_feature
<222> (532)..(532)

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<223> n is a, c, g, or t

<220>

<221> misc_feature

<222> (559)..(559)

<223> n is a, c, g, or t

<400> 136

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gagatattga ggggctgtaa gctaaagggt tgaaatctaa agatagagt gaaaaggagg	180
taggcacccc caccagcccc tgcttgacat ctgctttagt tcattccagc aggaaggagg	240
agaagggcag ggaggtaggg agctttctcag caaagccagc .ttcagctttg ataatctcac	300
ccacctaccc catttaagga gttccagggt taagagtta aaaacagggt gcaccagacc	360
atcattcagg agacaggaac tcattccagg ttctaagag aactcctatc tcagacctga	420
aggggtttcc anggcttcag nttgagctcc tctgggctaa ccagnagtca cttgattaag	480
tcccgctgcc ttgancccat nttccaggag ggggctattg cccgaggagg tntaaggagg	540
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<210> 137

<211> 1551

<212> DNA

<213> Homo sapiens

<400> 137

aaaacatggt gatcccaatg atgtgatcac tttgaacct ttccattaca aagcattgta	60
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attatgggta ctttaaagtc agtatttatc aagaaagga acttgaccac cattggcaca	180
tgtgacattt aagctcttca gccttttctt ttttagttgt aggtgtttac atttcatttc	240
taagccaact ctgtatttat gagagaagtt taagccttac atcatttgat actaaagggt	300
tatttggtgt aaatgaaaaa tgaccccaaa attacagagg aatatgccag ttttaagaaat	360
ggctacttaa agttgcttct ctctttcctt ctactcatg aaattaattg gtcttcttca	420
agtttcttta gattccatta aatgattaaa tcactattaa gagccattca tcaacgtgat	480
ttgtgtgtta gccaatgaat ctgtctcagc ttttgaccaa atgggtttta gacaaatgca	540
aagatctgcc tctagtcct atggctcttt ttgagtgcta gtattttgca tttcacataa	600
tgtagttatt ttgagctttt aaagagagca tttagacaaa gaagcaaaga gaggaagggg	660

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ccaatcaact catcagttcc atgcatcaac aaagcatagc tagtagagga atataaatga      720
cagattgaca aactgtagga aacactgtta ctctctttct gaagttttca agcaccatcc      780
tatgtgaaag ttccctcctg tccaaacaag ctcaaggccc atcttctccc tatacaaggc      840
aaacctgtaa ggccttcctt ccaaagagta cattgctttg gttttcttcc taaattccta      900
ttggaattag aactctcaga atccctggga gacagagcaa agatgactta attcattgag      960
cagcagagct ccctataagt gaacatcacc ttcccatctt ttccctactgc cacaccata    1020
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aattccttac cgaattttct cagatatacc tcatagacaa tagtgtttag agtaatgtta    1500
ttatagcgta tgtaataaat tattcactgt ttcttttggt aactgtgatt t              1551

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<210> 138
<211> 976
<212> PRT
<213> Homo sapiens

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<400> 138

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Met Glu Leu Gln Ala Ala Arg Ala Cys Phe Ala Leu Leu Trp Gly Cys
1      5      10      15

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Ala Leu Ala Ala Ala Ala Ala Ala Gln Gly Lys Glu Val Val Leu Leu
20      25      30

```

```

Asp Phe Ala Ala Ala Gly Gly Glu Leu Gly Trp Leu Thr His Pro Tyr
35      40      45

```

```

Gly Lys Gly Trp Asp Leu Met Gln Asn Ile Met Asn Asp Met Pro Ile
50      55      60

```

```

Tyr Met Tyr Ser Val Cys Asn Val Met Ser Gly Asp Gln Asp Asn Trp
65      70      75      80

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Leu Arg Thr Asn Trp Val Tyr Arg Gly Glu Ala Glu Arg Asn Asn Phe
85 90 95

Glu Leu Asn Phe Thr Val Arg Asp Cys Asn Ser Phe Pro Gly Gly Ala
100 105 110

Ser Ser Cys Lys Glu Thr Phe Asn Leu Tyr Tyr Ala Glu Ser Asp Leu
115 120 125

Asp Tyr Gly Thr Asn Phe Gln Lys Arg Leu Phe Thr Lys Ile Asp Thr
130 135 140

Ile Ala Pro Asp Glu Ile Thr Val Ser Ser Asp Phe Glu Ala Arg His
145 150 155 160

Val Lys Leu Asn Val Glu Glu Arg Ser Val Gly Pro Leu Thr Arg Lys
165 170 175

Gly Phe Tyr Leu Ala Phe Gln Asp Ile Gly Ala Cys Val Ala Leu Leu
180 185 190

Ser Val Arg Val Tyr Tyr Lys Lys Cys Pro Glu Leu Leu Gln Gly Leu
195 200 205

Ala His Phe Pro Glu Thr Ile Ala Gly Ser Asp Ala Pro Ser Leu Ala
210 215 220

Thr Val Ala Gly Thr Cys Val Asp His Ala Val Val Pro Pro Gly Gly
225 230 235 240

Glu Glu Pro Arg Met His Cys Ala Val Asp Gly Glu Trp Leu Val Pro
245 250 255

Ile Gly Gln Cys Leu Cys Gln Ala Gly Tyr Glu Lys Val Glu Asp Ala
260 265 270

Cys Gln Ala Cys Ser Pro Gly Phe Phe Lys Phe Glu Ala Ser Glu Ser
275 280 285

Pro Cys Leu Glu Cys Pro Glu His Thr Leu Pro Ser Pro Glu Gly Ala
290 295 300

Thr Ser Cys Glu Cys Glu Glu Gly Phe Phe Arg Ala Pro Gln Asp Pro
 305 310 315 320

Ala Ser Met Pro Cys Thr Arg Pro Pro Ser Ala Pro His Tyr Leu Thr
 325 330 335

Ala Val Gly Met Gly Ala Lys Val Glu Leu Arg Trp Thr Pro Pro Gln
 340 345 350

Asp Ser Gly Gly Arg Glu Asp Ile Val Tyr Ser Val Thr Cys Glu Gln
 355 360 365

Cys Trp Pro Glu Ser Gly Glu Cys Gly Pro Cys Glu Ala Ser Val Arg
 370 375 380

Tyr Ser Glu Pro Pro His Gly Leu Thr Arg Thr Ser Val Thr Val Ser
 385 390 395 400

Asp Leu Glu Pro His Met Asn Tyr Thr Phe Thr Val Glu Ala Arg Asn
 405 410 415

Gly Val Ser Gly Leu Val Thr Ser Arg Ser Phe Arg Thr Ala Ser Val
 420 425 430

Ser Ile Asn Gln Thr Glu Pro Pro Lys Val Arg Leu Glu Gly Arg Ser
 435 440 445

Thr Thr Ser Leu Ser Val Ser Trp Ser Ile Pro Pro Pro Gln Gln Ser
 450 455 460

Arg Val Trp Lys Tyr Glu Val Thr Tyr Arg Lys Lys Gly Asp Ser Asn
 465 470 475 480

Ser Tyr Asn Val Arg Arg Thr Glu Gly Phe Ser Val Thr Leu Asp Asp
 485 490 495

Leu Ala Pro Asp Thr Thr Tyr Leu Val Gln Val Gln Ala Leu Thr Gln
 500 505 510

Glu Gly Gln Gly Ala Gly Ser Lys Val His Glu Phe Gln Thr Leu Ser
 515 520 525

Pro Glu Gly Ser Gly Asn Leu Ala Val Ile Gly Gly Val Ala Val Gly

530		535		540
Val Val Leu Leu Leu Val Leu Ala Gly Val Gly Phe Phe Ile His Arg				
545		550	555	560
Arg Arg Lys Asn Gln Arg Ala Arg Gln Ser Pro Glu Asp Val Tyr Phe				
	565		570	575
Ser Lys Ser Glu Gln Leu Lys Pro Leu Lys Thr Tyr Val Asp Pro His				
	580		585	590
Thr Tyr Glu Asp Pro Asn Gln Ala Val Leu Lys Phe Thr Thr Glu Ile				
	595		600	605
His Pro Ser Cys Val Thr Arg Gln Lys Val Ile Gly Ala Gly Glu Phe				
	610		615	620
Gly Glu Val Tyr Lys Gly Met Leu Lys Thr Ser Ser Gly Lys Lys Glu				
	625		630	635
Val Pro Val Ala Ile Lys Thr Leu Lys Ala Gly Tyr Thr Glu Lys Gln				
	645		650	655
Arg Val Asp Phe Leu Gly Glu Ala Gly Ile Met Gly Gln Phe Ser His				
	660		665	670
His Asn Ile Ile Arg Leu Glu Gly Val Ile Ser Lys Tyr Lys Pro Met				
	675		680	685
Met Ile Ile Thr Glu Tyr Met Glu Asn Gly Ala Leu Asp Lys Phe Leu				
	690		695	700
Arg Glu Lys Asp Gly Glu Phe Ser Val Leu Gln Leu Val Gly Met Leu				
	705		710	715
Arg Gly Ile Ala Ala Gly Met Lys Tyr Leu Ala Asn Met Asn Tyr Val				
	725		730	735
His Arg Asp Leu Ala Ala Arg Asn Ile Leu Val Asn Ser Asn Leu Val				
	740		745	750
Cys Lys Val Ser Asp Phe Gly Leu Ser Arg Val Leu Glu Asp Asp Pro				
	755		760	765

Glu Ala Thr Tyr Thr Thr Ser Gly Gly Lys Ile Pro Ile Arg Trp Thr
 770 775 780

Ala Pro Glu Ala Ile Ser Tyr Arg Lys Phe Thr Ser Ala Ser Asp Val
 785 790 795 800

Trp Ser Phe Gly Ile Val Met Trp Glu Val Met Thr Tyr Gly Glu Arg
 805 810 815

Pro Tyr Trp Glu Leu Ser Asn His Glu Val Met Lys Ala Ile Asn Asp
 820 825 830

Gly Phe Arg Leu Pro Thr Pro Met Asp Cys Pro Ser Ala Ile Tyr Gln
 835 840 845

Leu Met Met Gln Cys Trp Gln Gln Glu Arg Ala Arg Arg Pro Lys Phe
 850 855 860

Ala Asp Ile Val Ser Ile Leu Asp Lys Leu Ile Arg Ala Pro Asp Ser
 865 870 875 880

Leu Lys Thr Leu Ala Asp Phe Asp Pro Arg Val Ser Ile Arg Leu Pro
 885 890 895

Ser Thr Ser Gly Ser Glu Gly Val Pro Phe Arg Thr Val Ser Glu Trp
 900 905 910

Leu Glu Ser Ile Lys Met Gln Gln Tyr Thr Glu His Phe Met Ala Ala
 915 920 925

Gly Tyr Thr Ala Ile Glu Lys Val Val Gln Met Thr Asn Asp Asp Ile
 930 935 940

Lys Arg Ile Gly Val Arg Leu Pro Gly His Gln Lys Arg Ile Ala Tyr
 945 950 955 960

Ser Leu Leu Gly Leu Lys Asp Gln Val Asn Thr Val Gly Ile Pro Ile
 965 970 975

<210> 139
 <211> 1055
 <212> PRT

<213> Homo sapiens

<400> 139

Met Ala Leu Arg Arg Leu Gly Ala Ala Leu Leu Leu Leu Pro Leu Leu
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Ala Ala Val Glu Glu Thr Leu Met Asp Ser Thr Thr Ala Thr Ala Glu
 20 25 30

Leu Gly Trp Met Val His Pro Pro Ser Gly Trp Glu Glu Val Ser Gly
 35 40 45

Tyr Asp Glu Asn Met Asn Thr Ile Arg Thr Tyr Gln Val Cys Asn Val
 50 55 60

Phe Glu Ser Ser Gln Asn Asn Trp Leu Arg Thr Lys Phe Ile Arg Arg
 65 70 75 80

Arg Gly Ala His Arg Ile His Val Glu Met Lys Phe Ser Val Arg Asp
 85 90 95

Cys Ser Ser Ile Pro Ser Val Pro Gly Ser Cys Lys Glu Thr Phe Asn
 100 105 110

Leu Tyr Tyr Tyr Glu Ala Asp Phe Asp Ser Ala Thr Lys Thr Phe Pro
 115 120 125

Asn Trp Met Glu Asn Pro Trp Val Lys Val Asp Thr Ile Ala Ala Asp
 130 135 140

Glu Ser Phe Ser Gln Val Asp Leu Gly Gly Arg Val Met Lys Ile Asn
 145 150 155 160

Thr Glu Val Arg Ser Phe Gly Pro Val Ser Arg Ser Gly Phe Tyr Leu
 165 170 175

Ala Phe Gln Asp Tyr Gly Gly Cys Met Ser Leu Ile Ala Val Arg Val
 180 185 190

Phe Tyr Arg Lys Cys Pro Arg Ile Ile Gln Asn Gly Ala Ile Phe Gln
 195 200 205

Glu Thr Leu Ser Gly Ala Glu Ser Thr Ser Leu Val Ala Ala Arg Gly

210	215	220
Ser Cys Ile Ala Asn Ala Glu Glu Val Asp Val Pro Ile Lys Leu Tyr		
225	230	235 240
Cys Asn Gly Asp Gly Glu Trp Leu Val Pro Ile Gly Arg Cys Met Cys		
	245	250 255
Lys Ala Gly Phe Glu Ala Val Glu Asn Gly Thr Val Cys Arg Gly Cys		
	260	265 270
Pro Ser Gly Thr Phe Lys Ala Asn Gln Gly Asp Glu Ala Cys Thr His		
	275	280 285
Cys Pro Ile Asn Ser Arg Thr Thr Ser Glu Gly Ala Thr Asn Cys Val		
	290	295 300
Cys Arg Asn Gly Tyr Tyr Arg Ala Asp Leu Asp Pro Leu Asp Met Pro		
305	310	315 320
Cys Thr Thr Ile Pro Ser Ala Pro Gln Ala Val Ile Ser Ser Val Asn		
	325	330 335
Glu Thr Ser Leu Met Leu Glu Trp Thr Pro Pro Arg Asp Ser Gly Gly		
	340	345 350
Arg Glu Asp Leu Val Tyr Asn Ile Ile Cys Lys Ser Cys Gly Ser Gly		
	355	360 365
Arg Gly Ala Cys Thr Arg Cys Gly Asp Asn Val Gln Tyr Ala Pro Arg		
	370	375 380
Gln Leu Gly Leu Thr Glu Pro Arg Ile Tyr Ile Ser Asp Leu Leu Ala		
385	390	395 400
His Thr Gln Tyr Thr Phe Glu Ile Gln Ala Val Asn Gly Val Thr Asp		
	405	410 415
Gln Ser Pro Phe Ser Pro Gln Phe Ala Ser Val Asn Ile Thr Thr Asn		
	420	425 430
Gln Ala Ala Pro Ser Ala Val Ser Ile Met His Gln Val Ser Arg Thr		
	435	440 445

Val Asp Ser Ile Thr Leu Ser Trp Ser Gln Pro Asp Gln Pro Asn Gly
 450 455 460

Val Ile Leu Asp Tyr Glu Leu Gln Tyr Tyr Glu Lys Glu Leu Ser Glu
 465 470 475 480

Tyr Asn Ala Thr Ala Ile Lys Ser Pro Thr Asn Thr Val Thr Val Gln
 485 490 495

Gly Leu Lys Ala Gly Ala Ile Tyr Val Phe Gln Val Arg Ala Arg Thr
 500 505 510

Val Ala Gly Tyr Gly Arg Tyr Ser Gly Lys Met Tyr Phe Gln Thr Met
 515 520 525

Thr Glu Ala Glu Tyr Gln Thr Ser Ile Gln Glu Lys Leu Pro Leu Ile
 530 535 540

Ile Gly Ser Ser Ala Ala Gly Leu Val Phe Leu Ile Ala Val Val Val
 545 550 555 560

Ile Ala Ile Val Cys Asn Arg Arg Gly Phe Glu Arg Ala Asp Ser Glu
 565 570 575

Tyr Thr Asp Lys Leu Gln His Tyr Thr Ser Gly His Met Thr Pro Gly
 580 585 590

Met Lys Ile Tyr Ile Asp Pro Phe Thr Tyr Glu Asp Pro Asn Glu Ala
 595 600 605

Val Arg Glu Phe Ala Lys Glu Ile Asp Ile Ser Cys Val Lys Ile Glu
 610 615 620

Gln Val Ile Gly Ala Gly Glu Phe Gly Glu Val Cys Ser Gly His Leu
 625 630 635 640

Lys Leu Pro Gly Lys Arg Glu Ile Phe Val Ala Ile Lys Thr Leu Lys
 645 650 655

Ser Gly Tyr Thr Glu Lys Gln Arg Arg Asp Phe Leu Ser Glu Ala Ser
 660 665 670

Ile Met Gly Gln Phe Asp His Pro Asn Val Ile His Leu Glu Gly Val
 675 680 685

Val Thr Lys Ser Thr Pro Val Met Ile Ile Thr Glu Phe Met Glu Asn
 690 695 700

Gly Ser Leu Asp Ser Phe Leu Arg Gln Asn Asp Gly Gln Phe Thr Val
 705 710 715 720

Ile Gln Leu Val Gly Met Leu Arg Gly Ile Ala Ala Gly Met Lys Tyr
 725 730 735

Leu Ala Asp Met Asn Tyr Val His Arg Asp Leu Ala Ala Arg Asn Ile
 740 745 750

Leu Val Asn Ser Asn Leu Val Cys Lys Val Ser Asp Phe Gly Leu Ser
 755 760 765

Arg Phe Leu Glu Asp Asp Thr Ser Asp Pro Thr Tyr Thr Ser Ala Leu
 770 775 780

Gly Gly Lys Ile Pro Ile Arg Trp Thr Ala Pro Glu Ala Ile Gln Tyr
 785 790 795 800

Arg Lys Phe Thr Ser Ala Ser Asp Val Trp Ser Tyr Gly Ile Val Met
 805 810 815

Trp Glu Val Met Ser Tyr Gly Glu Arg Pro Tyr Trp Asp Met Thr Asn
 820 825 830

Gln Asp Val Ile Asn Ala Ile Glu Gln Asp Tyr Arg Leu Pro Pro Pro
 835 840 845

Met Asp Cys Pro Ser Ala Leu His Gln Leu Met Leu Asp Cys Trp Gln
 850 855 860

Lys Asp Arg Asn His Arg Pro Lys Phe Gly Gln Ile Val Asn Thr Leu
 865 870 875 880

Asp Lys Met Ile Arg Asn Pro Asn Ser Leu Lys Ala Met Ala Pro Leu
 885 890 895

Ser Ser Gly Ile Asn Leu Pro Leu Leu Asp Arg Thr Ile Pro Asp Tyr
 900 905 910

Thr Ser Phe Asn Thr Val Asp Glu Trp Leu Glu Ala Ile Lys Met Gly
 915 920 925

Gln Tyr Lys Glu Ser Phe Ala Asn Ala Gly Phe Thr Ser Phe Asp Val
 930 935 940

Val Ser Gln Met Met Met Glu Asp Ile Leu Arg Val Gly Val Thr Leu
 945 950 955 960

Ala Gly His Gln Lys Lys Ile Leu Asn Ser Ile Gln Val Met Arg Ala
 965 970 975

Gln Met Asn Gln Ile Gln Ser Val Glu Gly Gln Pro Leu Ala Arg Arg
 980 985 990

Pro Arg Ala Thr Gly Arg Thr Lys Arg Cys Gln Pro Arg Asp Val Thr
 995 1000 1005

Lys Lys Thr Cys Asn Ser Asn Asp Gly Lys Lys Lys Gly Met Gly
 1010 1015 1020

Lys Lys Lys Thr Asp Pro Gly Arg Gly Arg Glu Ile Gln Gly Ile
 1025 1030 1035

Phe Phe Lys Glu Asp Ser His Lys Glu Ser Asn Asp Cys Ser Cys
 1040 1045 1050

Gly Gly
 1055

<210> 140
 <211> 178
 <212> PRT
 <213> Homo sapiens

<400> 140

Met Ser Gly Gly Lys Tyr Val Asp Ser Glu Gly His Leu Tyr Thr Val
 1 5 10 15

Pro Ile Arg Glu Gln Gly Asn Ile Tyr Lys Pro Asn Asn Lys Ala Met
 20 25 30

Ala Asp Glu Leu Ser Glu Lys Gln Val Tyr Asp Ala His Thr Lys Glu
 35 40 45

Ile Asp Leu Val Asn Arg Asp Pro Lys His Leu Asn Asp Asp Val Val
 50 55 60

Lys Ile Asp Phe Glu Asp Val Ile Ala Glu Pro Glu Gly Thr His Ser
 65 70 75 80

Phe Asp Gly Ile Trp Lys Ala Ser Phe Thr Thr Phe Thr Val Thr Lys
 85 90 95

Tyr Trp Phe Tyr Arg Leu Leu Ser Ala Leu Phe Gly Ile Pro Met Ala
 100 105 110

Leu Ile Trp Gly Ile Tyr Phe Ala Ile Leu Ser Phe Leu His Ile Trp
 115 120 125

Ala Val Val Pro Cys Ile Lys Ser Phe Leu Ile Glu Ile Gln Cys Ile
 130 135 140

Ser Arg Val Tyr Ser Ile Tyr Val His Thr Val Cys Asp Pro Leu Phe
 145 150 155 160

Glu Ala Val Gly Lys Ile Phe Ser Asn Val Arg Ile Asn Leu Gln Lys
 165 170 175

Glu Ile

<210> 141
 <211> 162
 <212> PRT
 <213> Homo sapiens

<400> 141

Met Gly Leu Glu Thr Glu Lys Ala Asp Val Gln Leu Phe Met Asp Asp
 1 5 10 15

Asp Ser Tyr Ser His His Ser Gly Leu Glu Tyr Ala Asp Pro Glu Lys
 20 25 30

Phe Ala Asp Ser Asp Gln Asp Arg Asp Pro His Arg Leu Asn Ser His
 35 40 45

Leu Lys Leu Gly Phe Glu Asp Val Ile Ala Glu Pro Val Thr Thr His
 50 55 60

Ser Phe Asp Lys Val Trp Ile Cys Ser His Ala Leu Phe Glu Ile Ser
 65 70 75 80

Lys Tyr Val Met Tyr Lys Phe Leu Thr Val Phe Leu Ala Ile Pro Leu
 85 90 95

Ala Phe Ile Ala Gly Ile Leu Phe Ala Thr Leu Ser Cys Leu His Ile
 100 105 110

Trp Ile Leu Met Pro Phe Val Lys Thr Cys Leu Met Val Leu Pro Ser
 115 120 125

Val Gln Thr Ile Trp Lys Ser Val Thr Asp Val Ile Ile Ala Pro Leu
 130 135 140

Cys Thr Ser Val Gly Arg Cys Phe Ser Ser Val Ser Leu Gln Leu Ser
 145 150 155 160

Gln Asp

<210> 142
 <211> 346
 <212> PRT
 <213> Homo sapiens

<400> 142

Met Ala Met Val Ser Glu Phe Leu Lys Gln Ala Trp Phe Ile Glu Asn
 1 5 10 15

Glu Glu Gln Glu Tyr Val Gln Thr Val Lys Ser Ser Lys Gly Gly Pro
 20 25 30

Gly Ser Ala Val Ser Pro Tyr Pro Thr Phe Asn Pro Ser Ser Asp Val
 35 40 45

Ala Ala Leu His Lys Ala Ile Met Val Lys Gly Val Asp Glu Ala Thr
 50 55 60

Ile Ile Asp Ile Leu Thr Lys Arg Asn Asn Ala Gln Arg Gln Gln Ile
65 70 75 80

Lys Ala Ala Tyr Leu Gln Glu Thr Gly Lys Pro Leu Asp Glu Thr Leu
85 90 95

Lys Lys Ala Leu Thr Gly His Leu Glu Glu Val Val Leu Ala Leu Leu
100 105 110

Lys Thr Pro Ala Gln Phe Asp Ala Asp Glu Leu Arg Ala Ala Met Lys
115 120 125

Gly Leu Gly Thr Asp Glu Asp Thr Leu Ile Glu Ile Leu Ala Ser Arg
130 135 140

Thr Asn Lys Glu Ile Arg Asp Ile Asn Arg Val Tyr Arg Glu Glu Leu
145 150 155 160

Lys Arg Asp Leu Ala Lys Asp Ile Thr Ser Asp Thr Ser Gly Asp Phe
165 170 175

Arg Asn Ala Leu Leu Ser Leu Ala Lys Gly Asp Arg Ser Glu Asp Phe
180 185 190

Gly Val Asn Glu Asp Leu Ala Asp Ser Asp Ala Arg Ala Leu Tyr Glu
195 200 205

Ala Gly Glu Arg Arg Lys Gly Thr Asp Val Asn Val Phe Asn Thr Ile
210 215 220

Leu Thr Thr Arg Ser Tyr Pro Gln Leu Arg Arg Val Phe Gln Lys Tyr
225 230 235 240

Thr Lys Tyr Ser Lys His Asp Met Asn Lys Val Leu Asp Leu Glu Leu
245 250 255

Lys Gly Asp Ile Glu Lys Cys Leu Thr Ala Ile Val Lys Cys Ala Thr
260 265 270

Ser Lys Pro Ala Phe Phe Ala Glu Lys Leu His Gln Ala Met Lys Gly
275 280 285

Val Gly Thr Arg His Lys Ala Leu Ile Arg Ile Met Val Ser Arg Ser
 290 295 300

Glu Ile Asp Met Asn Asp Ile Lys Ala Phe Tyr Gln Lys Met Tyr Gly
 305 310 315 320

Ile Ser Leu Cys Gln Ala Ile Leu Asp Glu Thr Lys Gly Asp Tyr Glu
 325 330 335

Lys Ile Leu Val Ala Leu Cys Gly Gly Asn
 340 345

<210> 143
 <211> 339
 <212> PRT
 <213> Homo sapiens

<400> 143

Met Ser Thr Val His Glu Ile Leu Cys Lys Leu Ser Leu Glu Gly Asp
 1 5 10 15

His Ser Thr Pro Pro Ser Ala Tyr Gly Ser Val Lys Ala Tyr Thr Asn
 20 25 30

Phe Asp Ala Glu Arg Asp Ala Leu Asn Ile Glu Thr Ala Ile Lys Thr
 35 40 45

Lys Gly Val Asp Glu Val Thr Ile Val Asn Ile Leu Thr Asn Arg Ser
 50 55 60

Asn Ala Gln Arg Gln Asp Ile Ala Phe Ala Tyr Gln Arg Arg Thr Lys
 65 70 75 80

Lys Glu Leu Ala Ser Ala Leu Lys Ser Ala Leu Ser Gly His Leu Glu
 85 90 95

Thr Val Ile Leu Gly Leu Leu Lys Thr Pro Ala Gln Tyr Asp Ala Ser
 100 105 110

Glu Leu Lys Ala Ser Met Lys Gly Leu Gly Thr Asp Glu Asp Ser Leu
 115 120 125

Ile Glu Ile Ile Cys Ser Arg Thr Asn Gln Glu Leu Gln Glu Ile Asn

130 135 140
 Arg Val Tyr Lys Glu Met Tyr Lys Thr Asp Leu Glu Lys Asp Ile Ile
 145 . 150 155 160
 Ser Asp Thr Ser Gly Asp Phe Arg Lys Leu Met Val Ala Leu Ala Lys
 165 170 175
 Gly Arg Arg Ala Glu Asp Gly Ser Val Ile Asp Tyr Glu Leu Ile Asp
 180 185 190
 Gln Asp Ala Arg Asp Leu Tyr Asp Ala Gly Val Lys Arg Lys Gly Thr
 195 200 205
 Asp Val Pro Lys Trp Ile Ser Ile Met Thr Glu Arg Ser Val Pro His
 210 215 220
 Leu Gln Lys Val Phe Asp Arg Tyr Lys Ser Tyr Ser Pro Tyr Asp Met
 225 230 235 240
 Leu Glu Ser Ile Arg Lys Glu Val Lys Gly Asp Leu Glu Asn Ala Phe
 245 250 255
 Leu Asn Leu Val Gln Cys Ile Gln Asn Lys Pro Leu Tyr Phe Ala Asp
 260 265 270
 Arg Leu Tyr Asp Ser Met Lys Gly Lys Gly Thr Arg Asp Lys Val Leu
 275 280 285
 Ile Arg Ile Met Val Ser Arg Ser Glu Val Asp Met Leu Lys Ile Arg
 290 295 300
 Ser Glu Phe Lys Arg Lys Tyr Gly Lys Ser Leu Tyr Tyr Tyr Ile Gln
 305 310 315 320
 Gln Asp Thr Lys Gly Asp Tyr Gln Lys Ala Leu Leu Tyr Leu Cys Gly
 325 330 335
 Gly Asp Asp

<210> 144
 <211> 372

<212> PRT

<213> Homo sapiens

<400> 144

Met Ala Thr Ser Pro Gln Lys Ser Pro Ser Val Pro Lys Ser Pro Thr
 1 5 10 15

Pro Lys Ser Pro Pro Ser Arg Lys Lys Asp Asp Ser Phe Leu Gly Lys
 20 25 30

Leu Gly Gly Thr Leu Ala Arg Arg Lys Lys Ala Lys Glu Val Ser Glu
 35 40 45

Leu Gln Glu Glu Gly Met Asn Ala Ile Asn Leu Pro Leu Ser Pro Ile
 50 55 60

Pro Phe Glu Leu Asp Pro Glu Asp Thr Met Leu Glu Glu Asn Glu Val
 65 70 75 80

Arg Thr Met Val Asp Pro Asn Ser Arg Ser Asp Pro Lys Leu Gln Glu
 85 90 95

Leu Met Lys Val Leu Ile Asp Trp Ile Asn Asp Val Leu Val Gly Glu
 100 105 110

Arg Ile Ile Val Lys Asp Leu Ala Glu Asp Leu Tyr Asp Gly Gln Val
 115 120 125

Leu Gln Lys Leu Phe Glu Lys Leu Glu Ser Glu Lys Leu Asn Val Ala
 130 135 140

Glu Val Thr Gln Ser Glu Ile Ala Gln Lys Gln Lys Leu Gln Thr Val
 145 150 155 160

Leu Glu Lys Ile Asn Glu Thr Leu Lys Leu Pro Pro Arg Ser Ile Lys
 165 170 175

Trp Asn Val Asp Ser Val His Ala Lys Ser Leu Val Ala Ile Leu His
 180 185 190

Leu Leu Val Ala Leu Ser Gln Tyr Phe Arg Ala Pro Ile Arg Leu Pro
 195 200 205

Asp His Val Ser Ile Gln Val Val Val Val Gln Lys Arg Glu Gly Ile
 210 215 220

Leu Gln Ser Arg Gln Ile Gln Glu Glu Ile Thr Gly Asn Thr Glu Ala
 225 230 235 240

Leu Ser Gly Arg His Glu Arg Asp Ala Phe Asp Thr Leu Phe Asp His
 245 250 255

Ala Pro Asp Lys Leu Asn Val Val Lys Lys Thr Leu Ile Thr Phe Val
 260 265 270

Asn Lys His Leu Asn Lys Leu Asn Leu Glu Val Thr Glu Leu Glu Thr
 275 280 285

Gln Phe Ala Asp Gly Val Tyr Leu Val Leu Leu Met Gly Leu Leu Glu
 290 295 300

Gly Tyr Phe Val Pro Leu His Ser Phe Phe Leu Thr Pro Asp Ser Phe
 305 310 315 320

Glu Gln Lys Val Leu Asn Val Ser Phe Ala Phe Glu Leu Met Gln Asp
 325 330 335

Gly Gly Leu Glu Lys Pro Lys Pro Arg Pro Glu Asp Ile Val Asn Cys
 340 345 350

Asp Leu Lys Ser Thr Leu Arg Val Leu Tyr Asn Leu Phe Thr Lys Tyr
 355 360 365

Arg Asn Val Glu
 370

<210> 145
 <211> 397
 <212> PRT
 <213> Homo sapiens

<400> 145

Met Arg Ser Pro Ser Ala Ala Trp Leu Leu Gly Ala Ala Ile Leu Leu
 1 5 10 15

Ala Ala Ser Leu Ser Cys Ser Gly Thr Ile Gln Gly Thr Ser Arg Ser
 20 25 30

Ser Lys Gly Arg Ser Leu Ile Gly Lys Val Asp Gly Thr Ser His Val
 35 40 45

Thr Gly Lys Gly Val Thr Val Glu Thr Val Phe Ser Val Asp Glu Phe
 50 55 60

Ser Ala Ser Val Leu Thr Gly Lys Leu Thr Thr Val Phe Leu Pro Ile
 65 70 75 80

Val Tyr Thr Ile Val Phe Val Val Gly Leu Pro Ser Asn Gly Met Ala
 85 90 95

Leu Trp Val Phe Leu Phe Arg Thr Lys Lys Lys His Pro Ala Val Ile
 100 105 110

Tyr Met Ala Asn Leu Ala Leu Ala Asp Leu Leu Ser Val Ile Trp Phe
 115 120 125

Pro Leu Lys Ile Ala Tyr His Ile His Gly Asn Asn Trp Ile Tyr Gly
 130 135 140

Glu Ala Leu Cys Asn Val Leu Ile Gly Phe Phe Tyr Gly Asn Met Tyr
 145 150 155 160

Cys Ser Ile Leu Phe Met Thr Cys Leu Ser Val Gln Arg Tyr Trp Val
 165 170 175

Ile Val Asn Pro Met Gly His Ser Arg Lys Lys Ala Asn Ile Ala Ile
 180 185 190

Gly Ile Ser Leu Ala Ile Trp Leu Leu Ile Leu Leu Val Thr Ile Pro
 195 200 205

Leu Tyr Val Val Lys Gln Thr Ile Phe Ile Pro Ala Leu Asn Ile Thr
 210 215 220

Thr Cys His Asp Val Leu Pro Glu Gln Leu Leu Val Gly Asp Met Phe
 225 230 235 240

Asn Tyr Phe Leu Ser Leu Ala Ile Gly Val Phe Leu Phe Pro Ala Phe
 245 250 255

Leu Thr Ala Ser Ala Tyr Val Leu Met Ile Arg Met Leu Arg Ser Ser
 260 265 270

Ala Met Asp Glu Asn Ser Glu Lys Lys Arg Lys Arg Ala Ile Lys Leu
 275 280 285

Ile Val Thr Val Leu Ala Met Tyr Leu Ile Cys Phe Thr Pro Ser Asn
 290 295 300

Leu Leu Leu Val Val His Tyr Phe Leu Ile Lys Ser Gln Gly Gln Ser
 305 310 315 320

His Val Tyr Ala Leu Tyr Ile Val Ala Leu Cys Leu Ser Thr Leu Asn
 325 330 335

Ser Cys Ile Asp Pro Phe Val Tyr Tyr Phe Val Ser His Asp Phe Arg
 340 345 350

Asp His Ala Lys Asn Ala Leu Leu Cys Arg Ser Val Arg Thr Val Lys
 355 360 365

Gln Met Gln Val Ser Leu Thr Ser Lys Lys His Ser Arg Lys Ser Ser
 370 375 380

Ser Tyr Ser Ser Ser Thr Thr Val Lys Thr Ser Tyr
 385 390 395

<210> 146
 <211> 295
 <212> PRT
 <213> Homo sapiens

<400> 146

Met Glu Thr Pro Ala Trp Pro Arg Val Pro Arg Pro Glu Thr Ala Val
 1 5 10 15

Ala Arg Thr Leu Leu Leu Gly Trp Val Phe Ala Gln Val Ala Gly Ala
 20 25 30

Ser Gly Thr Thr Asn Thr Val Ala Ala Tyr Asn Leu Thr Trp Lys Ser
 35 40 45

Thr Asn Phe Lys Thr Ile Leu Glu Trp Glu Pro Lys Pro Val Asn Gln

50		55		60
Val Tyr Thr	Val Gln Ile	Ser Thr Lys	Ser Gly Asp	Trp Lys Ser Lys
65	70		75	80
Cys Phe Tyr	Thr Thr Asp	Thr Glu Cys	Asp Leu Thr	Asp Glu Ile Val
	85		90	95
Lys Asp Val	Lys Gln Thr	Tyr Leu Ala	Arg Val Phe	Ser Tyr Pro Ala
	100		105	110
Gly Asn Val	Glu Ser Thr	Gly Ser Ala	Gly Glu Pro	Leu Tyr Glu Asn
	115		120	125
Ser Pro Glu	Phe Thr Pro	Tyr Leu Glu	Thr Asn Leu	Gly Gln Pro Thr
	130		135	140
Ile Gln Ser	Phe Glu Gln	Val Gly Thr	Lys Val Asn	Val Thr Val Glu
145		150		155
Asp Glu Arg	Thr Leu Val	Arg Arg Asn	Asn Thr Phe	Leu Ser Leu Arg
	165		170	175
Asp Val Phe	Gly Lys Asp	Leu Ile Tyr	Thr Leu Tyr	Tyr Trp Lys Ser
	180		185	190
Ser Ser Ser	Gly Lys Lys	Thr Ala Lys	Thr Asn Thr	Asn Glu Phe Leu
	195		200	205
Ile Asp Val	Asp Lys Gly	Glu Asn Tyr	Cys Phe Ser	Val Gln Ala Val
	210		215	220
Ile Pro Ser	Arg Thr Val	Asn Arg Lys	Ser Thr Asp	Ser Pro Val Glu
225		230		235
Cys Met Gly	Gln Glu Lys	Gly Glu Phe	Arg Glu Ile	Phe Tyr Ile Ile
	245		250	255
Gly Ala Val	Val Phe Val	Val Ile Ile	Leu Val Ile	Ile Ile Leu Ala Ile
	260		265	270
Ser Leu His	Lys Cys Arg	Lys Ala Gly	Val Gly Gln	Ser Trp Lys Glu
	275		280	285

Asn Ser Pro Leu Asn Val Ser
290 295

<210> 147
<211> 491
<212> PRT
<213> Homo sapiens

<400> 147

Met Ala Gln Ser Gly Gly Glu Ala Arg Pro Gly Pro Lys Thr Ala Val
1 5 10 15

Gln Ile Arg Val Ala Ile Gln Glu Ala Glu Asp Val Asp Glu Leu Glu
20 25 30

Asp Glu Glu Glu Gly Ala Glu Thr Arg Gly Ala Gly Asp Pro Ala Arg
35 40 45

Tyr Leu Ser Pro Gly Trp Gly Ser Ala Ser Glu Glu Glu Pro Ser Arg
50 55 60

Gly His Ser Gly Thr Thr Ala Ser Gly Gly Glu Asn Glu Arg Glu Asp
65 70 75 80

Leu Glu Gln Glu Trp Lys Pro Pro Asp Glu Glu Leu Ile Lys Lys Leu
85 90 95

Val Asp Gln Ile Glu Phe Cys Phe Ser Asp Glu Asn Leu Glu Lys Asp
100 105 110

Ala Phe Leu Leu Lys His Val Arg Arg Asn Lys Leu Gly Tyr Val Ser
115 120 125

Val Lys Leu Leu Thr Ser Phe Lys Lys Val Lys His Leu Thr Arg Asp
130 135 140

Trp Arg Thr Thr Ala His Ala Leu Lys Tyr Ser Val Val Leu Glu Leu
145 150 155 160

Asn Glu Asp His Arg Lys Val Arg Arg Thr Thr Pro Val Pro Leu Phe
165 170 175

Pro Asn Glu Asn Leu Pro Ser Lys Met Leu Leu Val Tyr Asp Leu Tyr
 180 185 190

Leu Ser Pro Lys Leu Trp Ala Leu Ala Thr Pro Gln Lys Asn Gly Arg
 195 200 205

Val Gln Glu Lys Val Met Glu His Leu Leu Lys Leu Phe Gly Thr Phe
 210 215 220

Gly Val Ile Ser Ser Val Arg Ile Leu Lys Pro Gly Arg Glu Leu Pro
 225 230 235 240

Pro Asp Ile Arg Arg Ile Ser Ser Arg Tyr Ser Gln Val Gly Thr Gln
 245 250 255

Glu Cys Ala Ile Val Glu Phe Glu Glu Val Glu Ala Ala Ile Lys Ala
 260 265 270

His Glu Phe Met Ile Thr Glu Ser Gln Gly Lys Glu Asn Met Lys Ala
 275 280 285

Val Leu Ile Gly Met Lys Pro Pro Lys Lys Lys Pro Ala Lys Asp Lys
 290 295 300

Asn His Asp Glu Glu Pro Thr Ala Ser Ile His Leu Asn Lys Ser Leu
 305 310 315 320

Asn Lys Arg Val Glu Glu Leu Gln Tyr Met Gly Asp Glu Ser Ser Ala
 325 330 335

Asn Ser Ser Ser Asp Pro Glu Ser Asn Pro Thr Ser Pro Met Ala Gly
 340 345 350

Arg Arg His Ala Ala Thr Asn Lys Leu Ser Pro Ser Gly His Gln Asn
 355 360 365

Leu Phe Leu Ser Pro Asn Ala Ser Pro Cys Thr Ser Pro Trp Ser Ser
 370 375 380

Pro Leu Ala Gln Arg Lys Gly Val Ser Arg Lys Ser Pro Leu Ala Glu
 385 390 395 400

Glu Gly Arg Leu Asn Cys Ser Thr Ser Pro Glu Ile Phe Arg Lys Cys

405 410 415
 Met Asp Tyr Ser Ser Asp Ser Ser Val Thr Pro Ser Gly Ser Pro Trp
 420 425 430
 Val Arg Arg Arg Arg Gln Ala Glu Met Gly Thr Gln Glu Lys Ser Pro
 435 440 445
 Gly Thr Ser Pro Leu Leu Ser Arg Lys Met Gln Thr Ala Asp Gly Leu
 450 455 460
 Pro Val Gly Val Leu Arg Leu Pro Arg Gly Pro Asp Asn Thr Arg Gly
 465 470 475 480
 Phe His Gly His Glu Arg Ser Arg Ala Cys Val
 485 490

 <210> 148
 <211> 374
 <212> PRT
 <213> Homo sapiens

 <400> 148
 Met Arg Pro Gly Thr Ala Leu Gln Ala Val Leu Leu Ala Val Leu Leu
 1 5 10 15
 Val Gly Leu Arg Ala Ala Thr Gly Arg Leu Leu Ser Gly Gln Pro Val
 20 25 30
 Cys Arg Gly Gly Thr Gln Arg Pro Cys Tyr Lys Val Ile Tyr Phe His
 35 40 45
 Asp Thr Ser Arg Arg Leu Asn Phe Glu Glu Ala Lys Glu Ala Cys Arg
 50 55 60
 Arg Asp Gly Gly Gln Leu Val Ser Ile Glu Ser Glu Asp Glu Gln Lys
 65 70 75 80
 Leu Ile Glu Lys Phe Ile Glu Asn Leu Leu Pro Ser Asp Gly Asp Phe
 85 90 95
 Trp Ile Gly Leu Arg Arg Arg Glu Glu Lys Gln Ser Asn Ser Thr Ala
 100 105 110

Cys Gln Asp Leu Tyr Ala Trp Thr Asp Gly Ser Ile Ser Gln Phe Arg
 115 120 125

Asn Trp Tyr Val Asp Glu Pro Ser Cys Gly Ser Glu Val Cys Val Val
 130 135 140

Met Tyr His Gln Pro Ser Ala Pro Ala Gly Ile Gly Gly Pro Tyr Met
 145 150 155 160

Phe Gln Trp Asn Asp Asp Arg Cys Asn Met Lys Asn Asn Phe Ile Cys
 165 170 175

Lys Tyr Ser Asp Glu Lys Pro Ala Val Pro Ser Arg Glu Ala Glu Gly
 180 185 190

Glu Glu Thr Glu Leu Thr Thr Pro Val Leu Pro Glu Glu Thr Gln Glu
 195 200 205

Glu Asp Ala Lys Lys Thr Phe Lys Glu Ser Arg Glu Ala Ala Leu Asn
 210 215 220

Leu Ala Tyr Ile Leu Ile Pro Ser Ile Pro Leu Leu Leu Leu Leu Val
 225 230 235 240

Val Thr Thr Val Val Cys Trp Val Trp Ile Cys Arg Lys Arg Lys Arg
 245 250 255

Glu Gln Pro Asp Pro Ser Thr Lys Lys Gln His Thr Ile Trp Pro Ser
 260 265 270

Pro His Gln Gly Asn Ser Pro Asp Leu Glu Val Tyr Asn Val Ile Arg
 275 280 285

Lys Gln Ser Glu Ala Asp Leu Ala Glu Thr Arg Pro Asp Leu Lys Asn
 290 295 300

Ile Ser Phe Arg Val Cys Ser Gly Glu Ala Thr Pro Asp Asp Met Ser
 305 310 315 320

Cys Asp Tyr Asp Asn Met Ala Val Asn Pro Ser Glu Ser Gly Phe Val
 325 330 335

Thr Leu Val Ser Val Glu Ser Gly Phe Val Thr Asn Asp Ile Tyr Glu
 340 345 350

Phe Ser Pro Asp Gln Met Gly Arg Ser Lys Glu Ser Gly Trp Val Glu
 355 360 365

Asn Glu Ile Tyr Gly Tyr
 370

<210> 149
 <211> 276
 <212> PRT
 <213> Homo sapiens

<400> 149

Met Ala Leu Leu Asp Val Cys Gly Ala Pro Arg Gly Gln Arg Pro Glu
 1 5 10 15

Ser Ala Leu Pro Val Ala Gly Ser Gly Arg Arg Ser Asp Pro Gly His
 20 25 30

Tyr Ser Phe Ser Met Arg Ser Pro Glu Leu Ala Leu Pro Arg Gly Met
 35 40 45

Lys Pro Thr Glu Phe Phe Gln Ser Leu Gly Gly Asp Gly Glu Arg Asn
 50 55 60

Val Gln Ile Glu Met Ala His Gly Thr Thr Thr Leu Ala Phe Lys Phe
 65 70 75 80

Gln His Gly Val Ile Ala Ala Val Asp Ser Arg Ala Ser Ala Gly Ser
 85 90 95

Tyr Ile Ser Ala Leu Arg Val Asn Lys Val Ile Glu Ile Asn Pro Tyr
 100 105 110

Leu Leu Gly Thr Met Ser Gly Cys Ala Ala Asp Cys Gln Tyr Trp Glu
 115 120 125

Arg Leu Leu Ala Lys Glu Cys Arg Leu Tyr Tyr Leu Arg Asn Gly Glu
 130 135 140

Arg Ile Ser Val Ser Ala Ala Ser Lys Leu Leu Ser Asn Met Met Cys
 145 150 155 160

Gln Tyr Arg Gly Met Gly Leu Ser Met Gly Ser Met Ile Cys Gly Trp
 165 170 175

Asp Lys Lys Gly Pro Gly Leu Tyr Tyr Val Asp Glu His Gly Thr Arg
 180 185 190

Leu Ser Gly Asn Met Phe Ser Thr Gly Ser Gly Asn Thr Tyr Ala Tyr
 195 200 205

Gly Val Met Asp Ser Gly Tyr Arg Pro Asn Leu Ser Pro Glu Glu Ala
 210 215 220

Tyr Asp Leu Gly Arg Arg Ala Ile Ala Tyr Ala Thr His Arg Asp Ser
 225 230 235 240

Tyr Ser Gly Gly Val Val Asn Met Tyr His Met Lys Glu Asp Gly Trp
 245 250 255

Val Lys Val Glu Ser Thr Asp Val Ser Asp Leu Leu His Gln Tyr Arg
 260 265 270

Glu Ala Asn Gln
 275

<210> 150
 <211> 219
 <212> PRT
 <213> Homo sapiens

<400> 150

Met Leu Arg Ala Gly Ala Pro Thr Gly Asp Leu Pro Arg Ala Gly Glu
 1 5 10 15

Val His Thr Gly Thr Thr Ile Met Ala Val Glu Phe Asp Gly Gly Val
 20 25 30

Val Met Gly Ser Asp Ser Arg Val Ser Ala Gly Glu Ala Val Val Asn
 35 40 45

Arg Val Phe Asp Lys Leu Ser Pro Leu His Glu Arg Ile Tyr Cys Ala
 50 55 60

Leu Ser Gly Ser Ala Ala Asp Ala Gln Ala Val Ala Asp Met Ala Ala
65 70 75 80

Tyr Gln Leu Glu Leu His Gly Ile Glu Leu Glu Glu Pro Pro Leu Val
85 90 95

Leu Ala Ala Ala Asn Val Val Arg Asn Ile Ser Tyr Lys Tyr Arg Glu
100 105 110

Asp Leu Ser Ala His Leu Met Val Ala Gly Trp Asp Gln Arg Glu Gly
115 120 125

Gly Gln Val Tyr Gly Thr Leu Gly Gly Met Leu Thr Arg Gln Pro Phe
130 135 140

Ala Ile Gly Gly Ser Gly Ser Thr Phe Ile Tyr Gly Tyr Val Asp Ala
145 150 155 160

Ala Tyr Lys Pro Gly Met Ser Pro Glu Glu Cys Arg Arg Phe Thr Thr
165 170 175

Asp Ala Ile Ala Leu Ala Met Ser Arg Asp Gly Ser Ser Gly Gly Val
180 185 190

Ile Tyr Leu Val Thr Ile Thr Ala Ala Gly Val Asp His Arg Val Ile
195 200 205

Leu Gly Asn Glu Leu Pro Lys Phe Tyr Asp Glu
210 215

<210> 151
<211> 253
<212> PRT
<213> Homo sapiens

<400> 151

Met Ala Asn Leu Gly Cys Trp Met Leu Val Leu Phe Val Ala Thr Trp
1 5 10 15

Ser Asp Leu Gly Leu Cys Lys Lys Arg Pro Lys Pro Gly Gly Trp Asn
20 25 30

Thr Gly Gly Ser Arg Tyr Pro Gly Gln Gly Ser Pro Gly Gly Asn Arg
35 40 45

Tyr Pro Pro Gln Gly Gly Gly Gly Trp Gly Gln Pro His Gly Gly Gly
50 55 60

Trp Gly Gln Pro His Gly Gly Gly Trp Gly Gln Pro His Gly Gly Gly
65 70 75 80

Trp Gly Gln Pro His Gly Gly Gly Trp Gly Gln Gly Gly Gly Thr His
85 90 95

Ser Gln Trp Asn Lys Pro Ser Lys Pro Lys Thr Asn Met Lys His Met
100 105 110

Ala Gly Ala Ala Ala Ala Gly Ala Val Val Gly Gly Leu Gly Gly Tyr
115 120 125

Met Leu Gly Ser Ala Met Ser Arg Pro Ile Ile His Phe Gly Ser Asp
130 135 140

Tyr Glu Asp Arg Tyr Tyr Arg Glu Asn Met His Arg Tyr Pro Asn Gln
145 150 155 160

Val Tyr Tyr Arg Pro Met Asp Glu Tyr Ser Asn Gln Asn Asn Phe Val
165 170 175

His Asp Cys Val Asn Ile Thr Ile Lys Gln His Thr Val Thr Thr Thr
180 185 190

Thr Lys Gly Glu Asn Phe Thr Glu Thr Asp Val Lys Met Met Glu Arg
195 200 205

Val Val Glu Gln Met Cys Ile Thr Gln Tyr Glu Arg Glu Ser Gln Ala
210 215 220

Tyr Tyr Gln Arg Gly Ser Ser Met Val Leu Phe Ser Ser Pro Pro Val
225 230 235 240

Ile Leu Leu Ile Ser Phe Leu Ile Phe Leu Ile Val Gly
245 250

<210> 152
<211> 323
<212> PRT

<213> Homo sapiens

<400> 152

Met Asp Ser Lys Gln Gln Cys Val Lys Leu Asn Asp Gly His Phe Met
 1 5 10 15

Pro Val Leu Gly Phe Gly Thr Tyr Ala Pro Pro Glu Val Pro Arg Ser
 20 25 30

Lys Ala Leu Glu Val Ser Lys Leu Ala Ile Glu Ala Gly Phe Arg His
 35 40 45

Ile Asp Ser Ala His Leu Tyr Asn Asn Glu Glu Gln Val Gly Leu Ala
 50 55 60

Ile Arg Ser Lys Ile Ala Asp Gly Ser Val Lys Arg Glu Asp Ile Phe
 65 70 75 80

Tyr Thr Ser Lys Leu Trp Ser Thr Ser His Arg Pro Glu Leu Val Arg
 85 90 95

Pro Ala Leu Glu Asn Ser Leu Lys Lys Ala Gln Leu Asp Tyr Val Asp
 100 105 110

Leu Tyr Leu Ile His Ser Pro Met Ser Leu Lys Pro Gly Glu Glu Leu
 115 120 125

Ser Pro Thr Asp Glu Asn Gly Lys Val Ile Phe Asp Ile Val Asp Leu
 130 135 140

Cys Thr Thr Trp Glu Ala Met Glu Lys Cys Lys Asp Ala Gly Leu Ala
 145 150 155 160

Lys Ser Ile Gly Val Ser Asn Phe Asn Arg Arg Gln Leu Glu Met Ile
 165 170 175

Leu Asn Lys Pro Gly Leu Lys Tyr Lys Pro Val Cys Asn Gln Val Glu
 180 185 190

Cys His Pro Tyr Phe Asn Arg Ser Lys Leu Leu Asp Phe Cys Lys Ser
 195 200 205

Lys Asp Ile Val Leu Val Ala Tyr Ser Ala Leu Gly Ser Gln Arg Asp

210 215 220
 Lys Arg Trp Val Asp Pro Asn Ser Pro Val Leu Leu Glu Asp Pro Val
 225 230 235 240
 Leu Cys Ala Leu Ala Lys Lys His Lys Arg Thr Pro Ala Leu Ile Ala
 245 250 255
 Leu Arg Tyr Gln Leu Gln Arg Gly Val Val Val Leu Ala Arg Ser Tyr
 260 265 270
 Asn Glu Gln Arg Ile Arg Gln Asn Val Gln Val Phe Glu Phe Gln Leu
 275 280 285
 Thr Ala Glu Asp Met Lys Ala Ile Asp Gly Leu Asp Arg Asn Leu His
 290 295 300
 Tyr Phe Asn Ser Asp Ser Phe Ala Ser His Pro Asn Tyr Pro Tyr Ser
 305 310 315 320
 Asp Glu Tyr

<210> 153
 <211> 784
 <212> PRT
 <213> Homo sapiens

<400> 153

Met Glu Gly Asp Gly Gly Thr Pro Trp Ala Leu Ala Leu Leu Arg Thr
 1 5 10 15
 Phe Asp Ala Gly Glu Phe Thr Gly Trp Glu Lys Val Gly Ser Gly Gly
 20 25 30
 Phe Gly Gln Val Tyr Lys Val Arg His Val His Trp Lys Thr Trp Leu
 35 40 45
 Ala Ile Lys Cys Ser Pro Ser Leu His Val Asp Asp Arg Glu Arg Met
 50 55 60
 Glu Leu Leu Glu Glu Ala Lys Lys Met Glu Met Ala Lys Phe Arg Tyr
 65 70 75 80

Ile Leu Pro Val Tyr Gly Ile Cys Arg Glu Pro Val Gly Leu Val Met
85 90 95

Glu Tyr Met Glu Thr Gly Ser Leu Glu Lys Leu Leu Ala Ser Glu Pro
100 105 110

Leu Pro Trp Asp Leu Arg Phe Arg Ile Ile His Glu Thr Ala Val Gly
115 120 125

Met Asn Phe Leu His Cys Met Ala Pro Pro Leu Leu His Leu Asp Leu
130 135 140

Lys Pro Ala Asn Ile Leu Leu Asp Ala His Tyr His Val Lys Ile Ser
145 150 155 160

Asp Phe Gly Leu Ala Lys Cys Asn Gly Leu Ser His Ser His Asp Leu
165 170 175

Ser Met Asp Gly Leu Phe Gly Thr Ile Ala Tyr Leu Pro Pro Glu Arg
180 185 190

Ile Arg Glu Lys Ser Arg Leu Phe Asp Thr Lys His Asp Val Tyr Ser
195 200 205

Phe Ala Ile Val Ile Trp Gly Val Leu Thr Gln Lys Lys Pro Phe Ala
210 215 220

Asp Glu Lys Asn Ile Leu His Ile Met Val Lys Val Val Lys Gly His
225 230 235 240

Arg Pro Glu Leu Pro Pro Val Cys Arg Ala Arg Pro Arg Ala Cys Ser
245 250 255

His Leu Ile Arg Leu Met Gln Arg Cys Trp Gln Gly Asp Pro Arg Val
260 265 270

Arg Pro Thr Phe Gln Glu Ile Thr Ser Glu Thr Glu Asp Leu Cys Glu
275 280 285

Lys Pro Asp Asp Glu Val Lys Glu Thr Ala His Asp Leu Asp Val Lys
290 295 300

Ser Pro Pro Glu Pro Arg Ser Glu Val Val Pro Ala Arg Leu Lys Arg
305 310 315 320

Ala Ser Ala Pro Thr Phe Asp Asn Asp Tyr Ser Leu Ser Glu Leu Leu
325 330 335

Ser Gln Leu Asp Ser Gly Val Ser Gln Ala Val Glu Gly Pro Glu Glu
340 345 350

Leu Ser Arg Ser Ser Ser Glu Ser Lys Leu Pro Ser Ser Gly Ser Gly
355 360 365

Lys Arg Leu Ser Gly Val Ser Ser Val Asp Ser Ala Phe Ser Ser Arg
370 375 380

Gly Ser Leu Ser Leu Ser Phe Glu Arg Glu Pro Ser Thr Ser Asp Leu
385 390 395 400

Gly Thr Thr Asp Val Gln Lys Lys Lys Leu Val Asp Ala Ile Val Ser
405 410 415

Gly Asp Thr Ser Lys Leu Met Lys Ile Leu Gln Pro Gln Asp Val Asp
420 425 430

Leu Ala Leu Asp Ser Gly Ala Ser Leu Leu His Leu Ala Val Glu Ala
435 440 445

Gly Gln Glu Glu Cys Ala Lys Trp Leu Leu Leu Asn Asn Ala Asn Pro
450 455 460

Asn Leu Ser Asn Arg Arg Gly Ser Thr Pro Leu His Met Ala Val Glu
465 470 475 480

Arg Arg Val Arg Gly Val Val Glu Leu Leu Leu Ala Arg Lys Ile Ser
485 490 495

Val Asn Ala Lys Asp Glu Asp Gln Trp Thr Ala Leu His Phe Ala Ala
500 505 510

Gln Asn Gly Asp Glu Ser Ser Thr Arg Leu Leu Leu Glu Lys Asn Ala
515 520 525

Ser Val Asn Glu Val Asp Phe Glu Gly Arg Thr Pro Met His Val Ala

530 535 540
 Cys Gln His Gly Gln Glu Asn Ile Val Arg Ile Leu Leu Arg Arg Gly
 545 550 555 560
 Val Asp Val Ser Leu Gln Gly Lys Asp Ala Trp Leu Pro Leu His Tyr
 565 570 575
 Ala Ala Trp Gln Gly His Leu Pro Ile Val Lys Leu Leu Ala Lys Gln
 580 585 590
 Pro Gly Val Ser Val Asn Ala Gln Thr Leu Asp Gly Arg Thr Pro Leu
 595 600 605
 His Leu Ala Ala Gln Arg Gly His Tyr Arg Val Ala Arg Ile Leu Ile
 610 615 620
 Asp Leu Cys Ser Asp Val Asn Val Cys Ser Leu Leu Ala Gln Thr Pro
 625 630 635 640
 Leu His Val Ala Ala Glu Thr Gly His Thr Ser Thr Ala Arg Leu Leu
 645 650 655
 Leu His Arg Gly Ala Gly Lys Lys Ala Val Thr Ser Asp Gly Tyr Thr
 660 665 670
 Ala Leu His Leu Ala Ala Arg Asn Gly His Leu Ala Thr Val Lys Leu
 675 680 685
 Leu Val Glu Glu Lys Ala Asp Val Leu Ala Arg Gly Pro Leu Asn Gln
 690 695 700
 Thr Ala Leu His Leu Ala Ala Ala His Gly His Ser Glu Val Val Glu
 705 710 715 720
 Glu Leu Val Ser Ala Asp Val Ile Asp Leu Phe Asp Glu Gln Gly Leu
 725 730 735
 Ser Ala Leu His Leu Ala Ala Gln Gly Arg His Ala Gln Thr Val Glu
 740 745 750
 Thr Leu Leu Arg His Gly Ala His Ile Asn Leu Gln Ser Leu Lys Phe
 755 760 765

Gln Gly Gly His Gly Pro Ala Ala Thr Leu Leu Arg Arg Ser Lys Thr
 770 775 780

<210> 154
 <211> 682
 <212> PRT
 <213> Homo sapiens

<400> 154

Met Gly Lys Lys Tyr Lys Asn Ile Val Leu Leu Lys Gly Leu Glu Val
 1 5 10 15

Ile Asn Asp Tyr His Phe Arg Met Val Lys Ser Leu Leu Ser Asn Asp
 20 25 30

Leu Lys Leu Asn Leu Lys Met Arg Glu Glu Tyr Asp Lys Ile Gln Ile
 35 40 45

Ala Asp Leu Met Glu Glu Lys Phe Arg Gly Asp Ala Gly Leu Gly Lys
 50 55 60

Leu Ile Lys Ile Phe Glu Asp Ile Pro Thr Leu Glu Asp Leu Ala Glu
 65 70 75 80

Thr Leu Lys Lys Glu Lys Leu Lys Val Lys Gly Pro Ala Leu Ser Arg
 85 90 95

Lys Arg Lys Lys Glu Val His Ala Thr Ser Pro Ala Pro Ser Thr Ser
 100 105 110

Ser Thr Val Lys Thr Glu Gly Ala Glu Ala Thr Pro Gly Ala Gln Lys
 115 120 125

Arg Lys Lys Ser Thr Lys Glu Lys Ala Gly Pro Lys Gly Ser Lys Val
 130 135 140

Ser Glu Glu Gln Thr Gln Pro Pro Ser Pro Ala Gly Ala Gly Met Ser
 145 150 155 160

Thr Ala Met Gly Arg Ser Pro Ser Pro Lys Thr Ser Leu Ser Ala Pro
 165 170 175

Pro Asn Ser Ser Ser Thr Glu Asn Pro Lys Thr Val Ala Lys Cys Gln
 180 185 190

Val Thr Pro Arg Arg Asn Val Leu Gln Lys Arg Pro Val Ile Val Lys
 195 200 205

Val Leu Ser Thr Thr Lys Pro Phe Glu Tyr Glu Thr Pro Glu Met Glu
 210 215 220

Lys Lys Ile Met Phe His Ala Thr Val Ala Thr Gln Thr Gln Phe Phe
 225 230 235 240

His Val Lys Val Leu Asn Thr Ser Leu Lys Glu Lys Phe Asn Gly Lys
 245 250 255

Lys Ile Ile Ile Ile Ser Asp Tyr Leu Glu Tyr Asp Ser Leu Leu Glu
 260 265 270

Val Asn Glu Glu Ser Thr Val Ser Glu Ala Gly Pro Asn Gln Thr Phe
 275 280 285

Glu Val Pro Asn Lys Ile Ile Asn Arg Ala Lys Glu Thr Leu Lys Ile
 290 295 300

Asp Ile Leu His Lys Gln Ala Ser Gly Asn Ile Val Tyr Gly Val Phe
 305 310 315 320

Met Leu His Lys Lys Thr Val Asn Gln Lys Thr Thr Ile Tyr Glu Ile
 325 330 335

Gln Asp Asp Arg Gly Lys Met Asp Val Val Gly Thr Gly Gln Cys His
 340 345 350

Asn Ile Pro Cys Glu Glu Gly Asp Lys Leu Gln Leu Phe Cys Phe Arg
 355 360 365

Leu Arg Lys Lys Asn Gln Met Ser Lys Leu Ile Ser Glu Met His Ser
 370 375 380

Phe Ile Gln Ile Lys Lys Lys Thr Asn Pro Arg Asn Asn Asp Pro Lys
 385 390 395 400

Ser Met Lys Leu Pro Gln Glu Gln Arg Gln Leu Pro Tyr Pro Ser Glu

405 410 415
 Ala Ser Thr Thr Phe Pro Glu Ser His Leu Arg Thr Pro Gln Met Pro
 420 425 430
 Pro Thr Thr Pro Ser Ser Ser Phe Phe Thr Lys Lys Ser Glu Asp Thr
 435 440 445
 Ile Ser Lys Met Asn Asp Phe Met Arg Met Gln Ile Leu Lys Glu Gly
 450 455 460
 Ser His Phe Pro Gly Pro Phe Met Thr Ser Ile Gly Pro Ala Glu Ser
 465 470 475 480
 His Pro His Thr Pro Gln Met Pro Pro Ser Thr Pro Ser Ser Ser Phe
 485 490 495
 Leu Thr Thr Leu Lys Pro Arg Leu Lys Thr Glu Pro Glu Glu Val Ser
 500 505 510
 Ile Glu Asp Ser Ala Gln Ser Asp Leu Lys Glu Val Met Val Leu Asn
 515 520 525
 Ala Thr Glu Ser Phe Val Tyr Glu Pro Lys Glu Gln Lys Lys Met Phe
 530 535 540
 His Ala Thr Val Ala Thr Glu Asn Glu Val Phe Arg Val Lys Val Phe
 545 550 555 560
 Asn Ile Asp Leu Lys Glu Lys Phe Thr Pro Lys Lys Ile Ile Ala Ile
 565 570 575
 Ala Asn Tyr Val Cys Arg Asn Gly Phe Leu Glu Val Tyr Pro Phe Thr
 580 585 590
 Leu Val Ala Asp Val Asn Ala Asp Arg Asn Met Glu Ile Pro Lys Gly
 595 600 605
 Leu Ile Arg Ser Ala Ser Val Thr Pro Lys Ile Asn Gln Leu Cys Ser
 610 615 620
 Gln Thr Lys Gly Ser Phe Val Asn Gly Val Phe Glu Val His Lys Val
 625 630 635 640

Ser Pro His His Cys Phe Ile Lys Phe Leu Leu Gln Pro Pro Ile Phe
 645 650 655

Lys Val Leu Thr Cys Gln Leu Glu Phe Gly Gln Leu Thr Gln His Arg
 660 665 670

Lys Ser Thr Pro Ser Pro Phe Pro Gln His
 675 680

<210> 155
 <211> 1218
 <212> PRT
 <213> Homo sapiens

<400> 155

Met Arg Ser Pro Arg Thr Arg Gly Arg Ser Gly Arg Pro Leu Ser Leu
 1 5 10 15

Leu Leu Ala Leu Leu Cys Ala Leu Arg Ala Lys Val Cys Gly Ala Ser
 20 25 30

Gly Gln Phe Glu Leu Glu Ile Leu Ser Met Gln Asn Val Asn Gly Glu
 35 40 45

Leu Gln Asn Gly Asn Cys Cys Gly Gly Ala Arg Asn Pro Gly Asp Arg
 50 55 60

Lys Cys Thr Arg Asp Glu Cys Asp Thr Tyr Phe Lys Val Cys Leu Lys
 65 70 75 80

Glu Tyr Gln Ser Arg Val Thr Ala Gly Gly Pro Cys Ser Phe Gly Ser
 85 90 95

Gly Ser Thr Pro Val Ile Gly Gly Asn Thr Phe Asn Leu Lys Ala Ser
 100 105 110

Arg Gly Asn Asp Arg Asn Arg Ile Val Leu Pro Phe Ser Phe Ala Trp
 115 120 125

Pro Arg Ser Tyr Thr Leu Leu Val Glu Ala Trp Asp Ser Ser Asn Asp
 130 135 140

Thr Val Gln Pro Asp Ser Ile Ile Glu Lys Ala Ser His Ser Gly Met
145 150 155 160

Ile Asn Pro Ser Arg Gln Trp Gln Thr Leu Lys Gln Asn Thr Gly Val
165 170 175

Ala His Phe Glu Tyr Gln Ile Arg Val Thr Cys Asp Asp Tyr Tyr Tyr
180 185 190

Gly Phe Gly Cys Asn Lys Phe Cys Arg Pro Arg Asp Asp Phe Phe Gly
195 200 205

His Tyr Ala Cys Asp Gln Asn Gly Asn Lys Thr Cys Met Glu Gly Trp
210 215 220

Met Gly Pro Glu Cys Asn Arg Ala Ile Cys Arg Gln Gly Cys Ser Pro
225 230 235 240

Lys His Gly Ser Cys Lys Leu Pro Gly Asp Cys Arg Cys Gln Tyr Gly
245 250 255

Trp Gln Gly Leu Tyr Cys Asp Lys Cys Ile Pro His Pro Gly Cys Val
260 265 270

His Gly Ile Cys Asn Glu Pro Trp Gln Cys Leu Cys Glu Thr Asn Trp
275 280 285

Gly Gly Gln Leu Cys Asp Lys Asp Leu Asn Tyr Cys Gly Thr His Gln
290 295 300

Pro Cys Leu Asn Gly Gly Thr Cys Ser Asn Thr Gly Pro Asp Lys Tyr
305 310 315 320

Gln Cys Ser Cys Pro Glu Gly Tyr Ser Gly Pro Asn Cys Glu Ile Ala
325 330 335

Glu His Ala Cys Leu Ser Asp Pro Cys His Asn Arg Gly Ser Cys Lys
340 345 350

Glu Thr Ser Leu Gly Phe Glu Cys Glu Cys Ser Pro Gly Trp Thr Gly
355 360 365

Pro Thr Cys Ser Thr Asn Ile Asp Asp Cys Ser Pro Asn Asn Cys Ser

370 375 380
 His Gly Gly Thr Cys Gln Asp Leu Val Asn Gly Phe Lys Cys Val Cys
 385 390 395 400
 Pro Pro Gln Trp Thr Gly Lys Thr Cys Gln Leu Asp Ala Asn Glu Cys
 405 410 415
 Glu Ala Lys Pro Cys Val Asn Ala Lys Ser Cys Lys Asn Leu Ile Ala
 420 425 430
 Ser Tyr Tyr Cys Asp Cys Leu Pro Gly Trp Met Gly Gln Asn Cys Asp
 435 440 445
 Ile Asn Ile Asn Asp Cys Leu Gly Gln Cys Gln Asn Asp Ala Ser Cys
 450 455 460
 Arg Asp Leu Val Asn Gly Tyr Arg Cys Ile Cys Pro Pro Gly Tyr Ala
 465 470 475 480
 Gly Asp His Cys Glu Arg Asp Ile Asp Glu Cys Ala Ser Asn Pro Cys
 485 490 495
 Leu Asn Gly Gly His Cys Gln Asn Glu Ile Asn Arg Phe Gln Cys Leu
 500 505 510
 Cys Pro Thr Gly Phe Ser Gly Asn Leu Cys Gln Leu Asp Ile Asp Tyr
 515 520 525
 Cys Glu Pro Asn Pro Cys Gln Asn Gly Ala Gln Cys Tyr Asn Arg Ala
 530 535 540
 Ser Asp Tyr Phe Cys Lys Cys Pro Glu Asp Tyr Glu Gly Lys Asn Cys
 545 550 555 560
 Ser His Leu Lys Asp His Cys Arg Thr Thr Pro Cys Glu Val Ile Asp
 565 570 575
 Ser Cys Thr Val Ala Met Ala Ser Asn Asp Thr Pro Glu Gly Val Arg
 580 585 590
 Tyr Ile Ser Ser Asn Val Cys Gly Pro His Gly Lys Cys Lys Ser Gln
 595 600 605

Ser Gly Gly Lys Phe Thr Cys Asp Cys Asn Lys Gly Phe Thr Gly Thr
610 615 620

Tyr Cys His Glu Asn Ile Asn Asp Cys Glu Ser Asn Pro Cys Arg Asn
625 630 635 640

Gly Gly Thr Cys Ile Asp Gly Val Asn Ser Tyr Lys Cys Ile Cys Ser
645 650 655

Asp Gly Trp Glu Gly Ala Tyr Cys Glu Thr Asn Ile Asn Asp Cys Ser
660 665 670

Gln Asn Pro Cys His Asn Gly Gly Thr Cys Arg Asp Leu Val Asn Asp
675 680 685

Phe Tyr Cys Asp Cys Lys Asn Gly Trp Lys Gly Lys Thr Cys His Ser
690 695 700

Arg Asp Ser Gln Cys Asp Glu Ala Thr Cys Asn Asn Gly Gly Thr Cys
705 710 715 720

Tyr Asp Glu Gly Asp Ala Phe Lys Cys Met Cys Pro Gly Gly Trp Glu
725 730 735

Gly Thr Thr Cys Asn Ile Ala Arg Asn Ser Ser Cys Leu Pro Asn Pro
740 745 750

Cys His Asn Gly Gly Thr Cys Val Val Asn Gly Glu Ser Phe Thr Cys
755 760 765

Val Cys Lys Glu Gly Trp Glu Gly Pro Ile Cys Ala Gln Asn Thr Asn
770 775 780

Asp Cys Ser Pro His Pro Cys Tyr Asn Ser Gly Thr Cys Val Asp Gly
785 790 795 800

Asp Asn Trp Tyr Arg Cys Glu Cys Ala Pro Gly Phe Ala Gly Pro Asp
805 810 815

Cys Arg Ile Asn Ile Asn Glu Cys Gln Ser Ser Pro Cys Ala Phe Gly
820 825 830

Ala Thr Cys Val Asp Glu Ile Asn Gly Tyr Arg Cys Val Cys Pro Pro
 835 840 845

Gly His Ser Gly Ala Lys Cys Gln Glu Val Ser Gly Arg Pro Cys Ile
 850 855 860

Thr Met Gly Ser Val Ile Pro Asp Gly Ala Lys Trp Asp Asp Asp Cys
 865 870 875 880

Asn Thr Cys Gln Cys Leu Asn Gly Arg Ile Ala Cys Ser Lys Val Trp
 885 890 895

Cys Gly Pro Arg Pro Cys Leu Leu His Lys Gly His Ser Glu Cys Pro
 900 905 910

Ser Gly Gln Ser Cys Ile Pro Ile Leu Asp Asp Gln Cys Phe Val His
 915 920 925

Pro Cys Thr Gly Val Gly Glu Cys Arg Ser Ser Ser Leu Gln Pro Val
 930 935 940

Lys Thr Lys Cys Thr Ser Asp Ser Tyr Tyr Gln Asp Asn Cys Ala Asn
 945 950 955 960

Ile Thr Phe Thr Phe Asn Lys Glu Met Met Ser Pro Gly Leu Thr Thr
 965 970 975

Glu His Ile Cys Ser Glu Leu Arg Asn Leu Asn Ile Leu Lys Asn Val
 980 985 990

Ser Ala Glu Tyr Ser Ile Tyr Ile Ala Cys Glu Pro Ser Pro Ser Ala
 995 1000 1005

Asn Asn Glu Ile His Val Ala Ile Ser Ala Glu Asp Ile Arg Asp
 1010 1015 1020

Asp Gly Asn Pro Ile Lys Glu Ile Thr Asp Lys Ile Ile Asp Leu
 1025 1030 1035

Val Ser Lys Arg Asp Gly Asn Ser Ser Leu Ile Ala Ala Val Ala
 1040 1045 1050

Glu Val Arg Val Gln Arg Arg Pro Leu Lys Asn Arg Thr Asp Phe
1055 1060 1065

Leu Val Pro Leu Leu Ser Ser Val Leu Thr Val Ala Trp Ile Cys
1070 1075 1080

Cys Leu Val Thr Ala Phe Tyr Trp Cys Leu Arg Lys Arg Arg Lys
1085 1090 1095

Pro Gly Ser His Thr His Ser Ala Ser Glu Asp Asn Thr Thr Asn
1100 1105 1110

Asn Val Arg Glu Gln Leu Asn Gln Ile Lys Asn Pro Ile Glu Lys
1115 1120 1125

His Gly Ala Asn Thr Val Pro Ile Lys Asp Tyr Glu Asn Lys Asn
1130 1135 1140

Ser Lys Met Ser Lys Ile Arg Thr His Asn Ser Glu Val Glu Glu
1145 1150 1155

Asp Asp Met Asp Lys His Gln Gln Lys Ala Arg Phe Ala Lys Gln
1160 1165 1170

Pro Ala Tyr Thr Leu Val Asp Arg Glu Glu Lys Pro Pro Asn Gly
1175 1180 1185

Thr Pro Thr Lys His Pro Asn Trp Thr Asn Lys Gln Asp Asn Arg
1190 1195 1200

Asp Leu Glu Ser Ala Gln Ser Leu Asn Arg Met Glu Tyr Ile Val
1205 1210 1215

<210> 156

<211> 334

<212> PRT

<213> Homo sapiens

<400> 156

Met Lys Met Ala Ser Ser Leu Ala Phe Leu Leu Leu Asn Phe His Val
1 5 10 15

Ser Leu Leu Leu Val Gln Leu Leu Thr Pro Cys Ser Ala Gln Phe Ser
20 25 30

Val Leu Gly Pro Ser Gly Pro Ile Leu Ala Met Val Gly Glu Asp Ala
35 40 45

Asp Leu Pro Cys His Leu Phe Pro Thr Met Ser Ala Glu Thr Met Glu
50 55 60

Leu Lys Trp Val Ser Ser Ser Leu Arg Gln Val Val Asn Val Tyr Ala
65 70 75 80

Asp Gly Lys Glu Val Glu Asp Arg Gln Ser Ala Pro Tyr Arg Gly Arg
85 90 95

Thr Ser Ile Leu Arg Asp Gly Ile Thr Ala Gly Lys Ala Ala Leu Arg
100 105 110

Ile His Asn Val Thr Ala Ser Asp Ser Gly Lys Tyr Leu Cys Tyr Phe
115 120 125

Gln Asp Gly Asp Phe Tyr Glu Lys Ala Leu Val Glu Leu Lys Val Ala
130 135 140

Ala Leu Gly Ser Asn Leu His Val Glu Val Lys Gly Tyr Glu Asp Gly
145 150 155 160

Gly Ile His Leu Glu Cys Arg Ser Thr Gly Trp Tyr Pro Gln Pro Gln
165 170 175

Ile Gln Trp Gly Asn Ala Lys Gly Glu Asn Ile Pro Ala Val Glu Ala
180 185 190

Pro Val Val Ala Asp Gly Val Gly Leu Tyr Glu Val Ala Ala Ser Val
195 200 205

Ile Met Lys Ser Gly Ser Gly Glu Gly Val Ser Cys Ile Ile Arg Asn
210 215 220

Ser Leu Leu Gly Leu Glu Lys Thr Ala Ser Ile Ser Ile Ala Asp Pro
225 230 235 240

Phe Phe Arg Ser Ala Gln Pro Trp Ile Ala Ala Leu Ala Gly Thr Leu
245 250 255

Pro Ile Leu Leu Leu Leu Leu Ala Gly Ala Ser Tyr Phe Leu Trp Arg
 260 265 270

Gln Gln Lys Glu Ile Thr Ala Leu Ser Ser Glu Ile Glu Ser Glu Gln
 275 280 285

Glu Met Lys Glu Met Gly Tyr Ala Ala Thr Glu Arg Glu Ile Ser Leu
 290 295 300

Arg Glu Ser Leu Gln Glu Glu Leu Lys Arg Lys Lys Ile Gln Tyr Leu
 305 310 315 320

Thr Arg Gly Glu Glu Ser Ser Ser Asp Thr Asn Lys Ser Ala
 325 330

<210> 157
 <211> 584
 <212> PRT
 <213> Homo sapiens

<400> 157

Met Lys Met Ala Ser Ser Leu Ala Phe Leu Leu Leu Asn Phe His Val
 1 5 10 15

Ser Leu Phe Leu Val Gln Leu Leu Thr Pro Cys Ser Ala Gln Phe Ser
 20 25 30

Val Leu Gly Pro Ser Gly Pro Ile Leu Ala Met Val Gly Glu Asp Ala
 35 40 45

Asp Leu Pro Cys His Leu Phe Pro Thr Met Ser Ala Glu Thr Met Glu
 50 55 60

Leu Arg Trp Val Ser Ser Ser Leu Arg Gln Val Val Asn Val Tyr Ala
 65 70 75 80

Asp Gly Lys Glu Val Glu Asp Arg Gln Ser Ala Pro Tyr Arg Gly Arg
 85 90 95

Thr Ser Ile Leu Arg Asp Gly Ile Thr Ala Gly Lys Ala Ala Leu Arg
 100 105 110

Ile His Asn Val Thr Ala Ser Asp Ser Gly Lys Tyr Leu Cys Tyr Phe

115 120 125
 Gln Asp Gly Asp Phe Tyr Glu Lys Ala Leu Val Glu Leu Lys Val Ala
 130 135 140
 Ala Leu Gly Ser Asp Leu His Ile Glu Val Lys Gly Tyr Glu Asp Gly
 145 150 155 160
 Gly Ile His Leu Glu Cys Arg Ser Thr Gly Trp Tyr Pro Gln Pro Gln
 165 170 175
 Ile Lys Trp Ser Asp Thr Lys Gly Glu Asn Ile Pro Ala Val Glu Ala
 180 185 190
 Pro Val Val Ala Asp Gly Val Gly Leu Tyr Ala Val Ala Ala Ser Val
 195 200 205
 Ile Met Arg Gly Ser Ser Gly Gly Gly Val Ser Cys Ile Ile Arg Asn
 210 215 220
 Ser Leu Leu Gly Leu Glu Lys Thr Ala Ser Ile Ser Ile Ala Asp Pro
 225 230 235 240
 Phe Phe Arg Ser Ala Gln Pro Trp Ile Ala Ala Leu Ala Gly Thr Leu
 245 250 255
 Pro Ile Ser Leu Leu Leu Leu Ala Gly Ala Ser Tyr Phe Leu Trp Arg
 260 265 270
 Gln Gln Lys Glu Lys Ile Ala Leu Ser Arg Glu Thr Glu Arg Glu Arg
 275 280 285
 Glu Met Lys Glu Met Gly Tyr Ala Ala Thr Glu Gln Glu Ile Ser Leu
 290 295 300
 Arg Glu Lys Leu Gln Glu Glu Leu Lys Trp Arg Lys Ile Gln Tyr Met
 305 310 315 320
 Ala Arg Gly Glu Lys Ser Leu Ala Tyr His Glu Trp Lys Met Ala Leu
 325 330 335
 Phe Lys Pro Ala Asp Val Ile Leu Asp Pro Asp Thr Ala Asn Ala Ile
 340 345 350

Leu Leu Val Ser Glu Asp Gln Arg Ser Val Gln Arg Ala Glu Glu Pro
 355 360 365

Arg Asp Leu Pro Asp Asn Pro Glu Arg Phe Glu Trp Arg Tyr Cys Val
 370 375 380

Leu Gly Cys Glu Asn Phe Thr Ser Gly Arg His Tyr Trp Glu Val Glu
 385 390 395 400

Val Gly Asp Arg Lys Glu Trp His Ile Gly Val Cys Ser Lys Asn Val
 405 410 415

Glu Arg Lys Lys Gly Trp Val Lys Met Thr Pro Glu Asn Gly Tyr Trp
 420 425 430

Thr Met Gly Leu Thr Asp Gly Asn Lys Tyr Arg Ala Leu Thr Glu Pro
 435 440 445

Arg Thr Asn Leu Lys Leu Pro Glu Pro Pro Arg Lys Val Gly Ile Phe
 450 455 460

Leu Asp Tyr Glu Thr Gly Glu Ile Ser Phe Tyr Asn Ala Thr Asp Gly
 465 470 475 480

Ser His Ile Tyr Thr Phe Pro His Ala Ser Phe Ser Glu Pro Leu Tyr
 485 490 495

Pro Val Phe Arg Ile Leu Thr Leu Glu Pro Thr Ala Leu Thr Ile Cys
 500 505 510

Pro Ile Pro Lys Glu Val Glu Ser Ser Pro Asp Pro Asp Leu Val Pro
 515 520 525

Asp His Ser Leu Glu Thr Pro Leu Thr Pro Gly Leu Ala Asn Glu Ser
 530 535 540

Gly Glu Pro Gln Ala Glu Val Thr Ser Leu Leu Leu Pro Ala His Pro
 545 550 555 560

Gly Ala Glu Val Ser Pro Ser Ala Thr Thr Asn Gln Asn His Lys Leu
 565 570 575

Gln Ala Arg Thr Glu Ala Leu Tyr
580

<210> 158
<211> 708
<212> PRT
<213> Homo sapiens

<400> 158

Met Asn Pro Thr Glu Thr Lys Ala Ile Pro Val Ser Gln Gln Met Glu
1 5 10 15

Gly Pro His Leu Pro Asn Lys Lys Lys His Lys Lys Gln Ala Val Lys
20 25 30

Thr Glu Pro Glu Lys Lys Ser Gln Ser Thr Lys Leu Ser Val Val His
35 40 45

Glu Lys Lys Ser Gln Glu Gly Lys Pro Lys Glu His Thr Glu Pro Lys
50 55 60

Ser Leu Pro Lys Gln Ala Ser Asp Thr Gly Ser Asn Asp Ala His Asn
65 70 75 80

Lys Lys Ala Val Ser Arg Ser Ala Glu Gln Gln Pro Ser Glu Lys Ser
85 90 95

Thr Glu Pro Lys Thr Lys Pro Gln Asp Met Ile Ser Ala Gly Gly Glu
100 105 110

Ser Val Ala Gly Ile Thr Ala Ile Ser Gly Lys Pro Gly Asp Lys Lys
115 120 125

Lys Glu Lys Lys Ser Leu Thr Pro Ala Val Pro Val Glu Ser Lys Pro
130 135 140

Asp Lys Pro Ser Gly Lys Ser Gly Met Asp Ala Ala Leu Asp Asp Leu
145 150 155 160

Ile Asp Thr Leu Gly Gly Pro Glu Glu Thr Glu Glu Glu Asn Thr Thr
165 170 175

Tyr Thr Gly Pro Glu Val Ser Asp Pro Met Ser Ser Thr Tyr Ile Glu

180	185	190
Glu Leu Gly Lys Arg Glu Val Thr Ile Pro Pro Lys Tyr Arg Glu Leu 195 200 205		
Leu Ala Lys Lys Glu Gly Ile Thr Gly Pro Pro Ala Asp Ser Ser Lys 210 215 220		
Pro Ile Gly Pro Asp Asp Ala Ile Asp Ala Leu Ser Ser Asp Phe Thr 225 230 235 240		
Cys Gly Ser Pro Thr Ala Ala Gly Lys Lys Thr Glu Lys Glu Glu Ser 245 250 255		
Thr Glu Val Leu Lys Ala Gln Ser Ala Gly Thr Val Arg Ser Ala Ala 260 265 270		
Pro Pro Gln Glu Lys Lys Arg Lys Val Glu Lys Asp Thr Met Ser Asp 275 280 285		
Gln Ala Leu Glu Ala Leu Ser Ala Ser Leu Gly Thr Arg Gln Ala Glu 290 295 300		
Pro Glu Leu Asp Leu Arg Ser Ile Lys Glu Val Asp Glu Ala Lys Ala 305 310 315 320		
Lys Glu Glu Lys Leu Glu Lys Cys Gly Glu Asp Asp Glu Thr Ile Pro 325 330 335		
Ser Glu Tyr Arg Leu Lys Pro Ala Thr Asp Lys Asp Gly Lys Pro Leu 340 345 350		
Leu Pro Glu Pro Glu Glu Lys Pro Lys Pro Arg Ser Glu Ser Glu Leu 355 360 365		
Ile Asp Glu Leu Ser Glu Asp Phe Asp Arg Ser Glu Cys Lys Glu Lys 370 375 380		
Pro Ser Lys Pro Thr Glu Lys Thr Glu Glu Ser Lys Ala Ala Ala Pro 385 390 395 400		
Ala Pro Val Ser Glu Ala Val Cys Arg Thr Ser Met Cys Ser Ile Gln 405 410 415		

Ser Ala Pro Pro Glu Pro Ala Thr Leu Lys Gly Thr Val Pro Asp Asp
 420 425 430

Ala Val Glu Ala Leu Ala Asp Ser Leu Gly Lys Lys Glu Ala Asp Pro
 435 440 445

Glu Asp Gly Lys Pro Val Met Asp Lys Val Lys Glu Lys Ala Lys Glu
 450 455 460

Glu Asp Arg Glu Lys Leu Gly Glu Lys Glu Glu Thr Ile Pro Pro Asp
 465 470 475 480

Tyr Arg Leu Glu Glu Val Lys Asp Lys Asp Gly Lys Pro Leu Leu Pro
 485 490 495

Lys Glu Ser Lys Glu Gln Leu Pro Pro Met Ser Glu Asp Phe Leu Leu
 500 505 510

Asp Ala Leu Ser Glu Asp Phe Ser Gly Pro Gln Asn Ala Ser Ser Leu
 515 520 525

Lys Phe Glu Asp Ala Lys Leu Ala Ala Ala Ile Ser Glu Val Val Ser
 530 535 540

Gln Thr Pro Ala Ser Thr Thr Gln Ala Gly Ala Pro Pro Arg Asp Thr
 545 550 555 560

Ser Gln Ser Asp Lys Asp Leu Asp Asp Ala Leu Asp Lys Leu Ser Asp
 565 570 575

Ser Leu Gly Gln Arg Gln Pro Asp Pro Asp Glu Asn Lys Pro Met Glu
 580 585 590

Asp Lys Val Lys Glu Lys Ala Lys Ala Glu His Arg Asp Lys Leu Gly
 595 600 605

Glu Arg Asp Asp Thr Ile Pro Pro Glu Tyr Arg His Leu Leu Asp Asp
 610 615 620

Asn Gly Gln Asp Lys Pro Val Lys Pro Pro Thr Lys Lys Ser Glu Asp
 625 630 635 640

Ser Lys Lys Pro Ala Asp Asp Gln Asp Pro Ile Asp Ala Leu Ser Gly
645 650 655

Asp Leu Asp Ser Cys Pro Ser Thr Thr Glu Thr Ser Gln Asn Thr Ala
660 665 670

Lys Asp Lys Cys Lys Lys Ala Ala Ser Ser Ser Lys Ala Pro Lys Asn
675 680 685

Gly Gly Lys Ala Lys Asp Ser Ala Lys Thr Thr Glu Glu Thr Ser Lys
690 695 700

Pro Lys Asp Asp
705

<210> 159
<211> 395
<212> PRT
<213> Homo sapiens

<400> 159

Met Trp Leu Pro Arg Val Ser Ser Thr Ala Val Thr Ala Leu Leu Leu
1 5 10 15

Ala Gln Thr Phe Leu Leu Leu Phe Leu Val Ser Arg Pro Gly Pro Ser
20 25 30

Ser Pro Ala Gly Gly Glu Ala Arg Val His Val Leu Val Leu Ser Ser
35 40 45

Trp Arg Ser Gly Ser Ser Phe Val Gly Gln Leu Phe Asn Gln His Pro
50 55 60

Asp Val Phe Tyr Leu Met Glu Pro Ala Trp His Val Trp Thr Thr Leu
65 70 75 80

Ser Gln Gly Ser Ala Ala Thr Leu His Met Ala Val Arg Asp Leu Val
85 90 95

Arg Ser Val Phe Leu Cys Asp Met Asp Val Phe Asp Ala Tyr Leu Pro
100 105 110

Trp Arg Arg Asn Leu Ser Asp Leu Phe Gln Trp Ala Val Ser Arg Ala

115	120	125
Leu Cys Ser Pro Pro Ala Cys Ser Ala Phe Pro Arg Gly Ala Ile Ser		
130	135	140
Ser Glu Ala Val Cys Lys Pro Leu Cys Ala Arg Gln Ser Phe Thr Leu		
145	150	155
Ala Arg Glu Ala Cys Arg Ser Tyr Ser His Val Val Leu Lys Glu Val		
165	170	175
Arg Phe Phe Asn Leu Gln Val Leu Tyr Pro Leu Leu Ser Asp Pro Ala		
180	185	190
Leu Asn Leu Arg Ile Val His Leu Val Arg Asp Pro Arg Ala Val Leu		
195	200	205
Arg Ser Arg Glu Gln Thr Ala Lys Ala Leu Ala Arg Asp Asn Gly Ile		
210	215	220
Val Leu Gly Thr Asn Gly Thr Trp Val Glu Ala Asp Pro Gly Leu Arg		
225	230	235
Val Val Arg Glu Val Cys Arg Ser His Val Arg Ile Ala Glu Ala Ala		
245	250	255
Thr Leu Lys Pro Pro Pro Phe Leu Arg Gly Arg Tyr Arg Leu Val Arg		
260	265	270
Phe Glu Asp Leu Ala Arg Glu Pro Leu Ala Glu Ile Arg Ala Leu Tyr		
275	280	285
Ala Phe Thr Gly Leu Ser Leu Thr Pro Gln Leu Glu Ala Trp Ile His		
290	295	300
Asn Ile Thr His Gly Ser Gly Pro Gly Ala Arg Arg Glu Ala Phe Lys		
305	310	315
Thr Ser Ser Arg Asn Ala Leu Asn Val Ser Gln Ala Trp Arg His Ala		
325	330	335
Leu Pro Phe Ala Lys Ile Arg Arg Val Gln Glu Leu Cys Ala Gly Ala		
340	345	350

Leu Gln Leu Leu Gly Tyr Arg Pro Val Tyr Ser Glu Asp Glu Gln Arg
 355 360 365

Asn Leu Ala Leu Asp Leu Val Leu Pro Arg Gly Leu Asn Gly Phe Thr
 370 375 380

Trp Ala Ser Ser Thr Ala Ser His Pro Arg Asn
 385 390 395

<210> 160
 <211> 254
 <212> PRT
 <213> Homo sapiens

<400> 160

Met Pro Ala Lys Thr Pro Ile Tyr Leu Lys Ala Ala Asn Asn Lys Lys
 1 5 10 15

Gly Lys Lys Phe Lys Leu Arg Asp Ile Leu Ser Pro Asp Met Ile Ser
 20 25 30

Pro Pro Leu Gly Asp Phe Arg His Thr Ile His Ile Gly Lys Glu Gly
 35 40 45

Gln His Asp Val Phe Gly Asp Ile Ser Phe Leu Gln Gly Asn Tyr Glu
 50 55 60

Leu Leu Pro Gly Asn Gln Glu Lys Ala His Leu Gly Gln Phe Pro Gly
 65 70 75 80

His Asn Glu Phe Phe Arg Ala Asn Ser Thr Ser Asp Ser Val Phe Thr
 85 90 95

Glu Thr Pro Ser Pro Val Leu Lys Asn Ala Ile Ser Leu Pro Thr Ile
 100 105 110

Gly Gly Ser Gln Ala Leu Met Leu Pro Leu Leu Ser Pro Val Thr Phe
 115 120 125

Asn Ser Lys Gln Glu Ser Phe Gly Pro Ala Lys Leu Pro Arg Leu Ser
 130 135 140

Cys Glu Pro Val Met Glu Glu Lys Ala Gln Glu Lys Ser Ser Leu Leu
 145 150 155 160

Glu Asn Gly Thr Val His Gln Gly Asp Thr Ser Trp Gly Ser Ser Gly
 165 170 175

Ser Ala Ser Gln Ser Ser Gln Gly Arg Asp Ser His Ser Ser Ser Leu
 180 185 190

Ser Glu Gln Tyr Pro Asp Trp Pro Ala Glu Asp Met Phe Asp His Pro
 195 200 205

Thr Pro Cys Glu Leu Ile Lys Gly Lys Thr Lys Ser Glu Glu Ser Leu
 210 215 220

Ser Asp Leu Thr Gly Ser Leu Leu Ser Leu Gln Leu Asp Leu Gly Pro
 225 230 235 240

Ser Leu Leu Asp Glu Val Leu Asn Val Met Asp Lys Asn Lys
 245 250

<210> 161
 <211> 536
 <212> PRT
 <213> Homo sapiens

<400> 161

Met Asp Lys Val Gly Lys Met Trp Asn Asn Phe Lys Tyr Arg Cys Gln
 1 5 10 15

Asn Leu Phe Gly His Glu Gly Gly Ser Arg Ser Glu Asn Val Asp Met
 20 25 30

Asn Ser Asn Arg Cys Leu Ser Val Lys Glu Lys Asn Ile Ser Ile Gly
 35 40 45

Asp Ser Thr Pro Gln Gln Gln Ser Ser Pro Leu Arg Glu Asn Ile Ala
 50 55 60

Leu Gln Leu Gly Leu Ser Pro Ser Lys Asn Ser Ser Arg Arg Asn Gln
 65 70 75 80

Asn Cys Ala Thr Glu Ile Pro Gln Ile Val Glu Ile Ser Ile Glu Lys
 85 90 95

Asp Asn Asp Ser Cys Val Thr Pro Gly Thr Arg Leu Ala Arg Arg Asp
 100 105 110

Ser Tyr Ser Arg His Ala Pro Trp Gly Gly Lys Lys Lys His Ser Cys
 115 120 125

Ser Thr Lys Thr Gln Ser Ser Leu Asp Ala Asp Lys Lys Phe Gly Arg
 130 135 140

Thr Arg Ser Gly Leu Gln Arg Arg Glu Arg Arg Tyr Gly Val Ser Ser
 145 150 155 160

Val His Asp Met Asp Ser Val Ser Ser Arg Thr Val Gly Ser Arg Ser
 165 170 175

Leu Arg Gln Arg Leu Gln Asp Thr Val Gly Leu Cys Phe Pro Met Arg
 180 185 190

Thr Tyr Ser Lys Gln Ser Lys Pro Leu Phe Ser Asn Lys Arg Lys Ile
 195 200 205

His Leu Ser Glu Leu Met Leu Glu Lys Cys Pro Phe Pro Ala Gly Ser
 210 215 220

Asp Leu Ala Gln Lys Trp His Leu Ile Lys Gln His Thr Ala Pro Val
 225 230 235 240

Ser Pro His Ser Thr Phe Phe Asp Thr Phe Asp Pro Ser Leu Val Ser
 245 250 255

Thr Glu Asp Glu Glu Asp Arg Leu Arg Glu Arg Arg Arg Leu Ser Ile
 260 265 270

Glu Glu Gly Val Asp Pro Pro Pro Asn Ala Gln Ile His Thr Phe Glu
 275 280 285

Ala Thr Ala Gln Val Asn Pro Leu Tyr Lys Leu Gly Pro Lys Leu Ala
 290 295 300

Pro Gly Met Thr Glu Ile Ser Gly Asp Ser Ser Ala Ile Pro Gln Ala
 305 310 315 320

Asn Cys Asp Ser Glu Glu Asp Thr Thr Thr Leu Cys Leu Gln Ser Arg
 325 330 335

Arg Gln Lys Gln Arg Gln Ile Ser Gly Asp Ser His Thr His Val Ser
 340 345 350

Arg Gln Gly Ala Trp Lys Val His Thr Gln Ile Asp Tyr Ile His Cys
 355 360 365

Leu Val Pro Asp Leu Leu Gln Ile Thr Gly Asn Pro Cys Tyr Trp Gly
 370 375 380

Val Met Asp Arg Tyr Glu Ala Glu Ala Leu Leu Glu Gly Lys Pro Glu
 385 390 395 400

Gly Thr Phe Leu Leu Arg Asp Ser Ala Gln Glu Asp Tyr Leu Phe Ser
 405 410 415

Val Ser Phe Arg Arg Tyr Asn Arg Ser Leu His Ala Arg Ile Glu Gln
 420 425 430

Trp Asn His Asn Phe Ser Phe Asp Ala His Asp Pro Cys Val Phe His
 435 440 445

Ser Ser Thr Val Thr Gly Leu Leu Glu His Tyr Lys Asp Pro Ser Ser
 450 455 460

Cys Met Phe Phe Glu Pro Leu Leu Thr Ile Ser Leu Asn Arg Thr Phe
 465 470 475 480

Pro Phe Ser Leu Gln Tyr Ile Cys Arg Ala Val Ile Cys Arg Cys Thr
 485 490 495

Thr Tyr Asp Gly Ile Asp Gly Leu Pro Leu Pro Ser Met Leu Gln Asp
 500 505 510

Phe Leu Lys Glu Tyr His Tyr Lys Gln Lys Val Arg Val Arg Trp Leu
 515 520 525

Glu Arg Glu Pro Val Lys Ala Lys
 530 535

<210> 162
 <211> 142
 <212> PRT
 <213> Homo sapiens

<400> 162

Met Ala Thr Lys Ile Asp Lys Glu Ala Cys Arg Ala Ala Tyr Asn Leu
 1 5 10 15

Val Arg Asp Asp Gly Ser Ala Val Ile Trp Val Thr Phe Lys Tyr Asp
 20 25 30

Gly Ser Thr Ile Val Pro Gly Glu Gln Gly Ala Glu Tyr Gln His Phe
 35 40 45

Ile Gln Gln Cys Thr Asp Asp Val Arg Leu Phe Ala Phe Val Arg Phe
 50 55 60

Thr Thr Gly Asp Ala Met Ser Lys Arg Ser Lys Phe Ala Leu Ile Thr
 65 70 75 80

Trp Ile Gly Glu Asn Val Ser Gly Leu Gln Arg Ala Lys Thr Gly Thr
 85 90 95

Asp Lys Thr Leu Val Lys Glu Val Val Gln Asn Phe Ala Lys Glu Phe
 100 105 110

Val Ile Ser Asp Arg Lys Glu Leu Glu Glu Asp Phe Ile Lys Ser Glu
 115 120 125

Leu Lys Lys Ala Gly Gly Ala Asn Tyr Asp Ala Gln Thr Glu
 130 135 140

<210> 163
 <211> 658
 <212> PRT
 <213> Homo sapiens

<400> 163

Met Ala Glu Ala Ala Ala Ala Ala Gly Gly Thr Gly Leu Gly Ala Gly
 1 5 10 15

Ala Ser Tyr Gly Ser Ala Ala Asp Arg Asp Arg Asp Pro Asp Pro Asp
 20 25 30

Arg Ala Gly Arg Arg Leu Arg Val Leu Ser Gly His Leu Leu Gly Arg
 35 40 45
 Pro Arg Glu Ala Leu Ser Thr Asn Glu Cys Lys Ala Arg Arg Ala Ala
 50 55 60
 Ser Ala Ala Thr Ala Ala Pro Thr Ala Thr Pro Ala Ala Gln Glu Ser
 65 70 75 80
 Gly Thr Ile Pro Lys Lys Arg Gln Glu Val Met Lys Trp Asn Gly Trp
 85 90 95
 Gly Tyr Asn Asp Ser Lys Phe Ile Phe Asn Lys Lys Gly Gln Ile Glu
 100 105 110
 Leu Thr Gly Lys Arg Tyr Pro Leu Ser Gly Met Gly Leu Pro Thr Phe
 115 120 125
 Lys Glu Trp Ile Gln Asn Thr Leu Gly Val Asn Val Glu His Lys Thr
 130 135 140
 Thr Ser Lys Ala Ser Leu Asn Pro Ser Asp Thr Pro Pro Ser Val Val
 145 150 155 160
 Asn Glu Asp Phe Leu His Asp Leu Lys Glu Thr Asn Ile Ser Tyr Ser
 165 170 175
 Gln Glu Ala Asp Asp Arg Val Phe Arg Ala His Gly His Cys Leu His
 180 185 190
 Glu Ile Phe Leu Leu Arg Glu Gly Met Phe Glu Arg Ile Pro Asp Ile
 195 200 205
 Val Leu Trp Pro Thr Cys His Asp Asp Val Val Lys Ile Val Asn Leu
 210 215 220
 Ala Cys Lys Tyr Asn Leu Cys Ile Ile Pro Ile Gly Gly Gly Thr Ser
 225 230 235 240
 Val Ser Tyr Gly Leu Met Cys Pro Ala Asp Glu Thr Arg Thr Ile Ile
 245 250 255

Ser Leu Asp Thr Ser Gln Met Asn Arg Ile Leu Trp Val Asp Glu Asn
 260 265 270

Asn Leu Thr Ala His Val Glu Ala Gly Ile Thr Gly Gln Glu Leu Glu
 275 280 285

Arg Gln Leu Lys Glu Ser Gly Tyr Cys Thr Gly His Glu Pro Asp Ser
 290 295 300

Leu Glu Phe Ser Thr Val Gly Gly Trp Val Ser Thr Arg Ala Ser Gly
 305 310 315 320

Met Lys Lys Asn Ile Tyr Gly Asn Ile Glu Asp Leu Val Val His Ile
 325 330 335

Lys Met Val Thr Pro Arg Gly Ile Ile Glu Lys Ser Cys Gln Gly Pro
 340 345 350

Arg Met Ser Thr Gly Pro Asp Ile His His Phe Ile Met Gly Ser Glu
 355 360 365

Gly Thr Leu Gly Val Ile Thr Glu Ala Thr Ile Lys Ile Arg Pro Val
 370 375 380

Pro Glu Tyr Gln Lys Tyr Gly Ser Val Ala Phe Pro Asn Phe Glu Gln
 385 390 395 400

Gly Val Ala Cys Leu Arg Glu Ile Ala Lys Gln Arg Cys Ala Pro Ala
 405 410 415

Ser Ile Arg Leu Met Asp Asn Lys Gln Phe Gln Phe Gly His Ala Leu
 420 425 430

Lys Pro Gln Val Ser Ser Ile Phe Thr Ser Phe Leu Asp Gly Leu Lys
 435 440 445

Lys Phe Tyr Ile Thr Lys Phe Lys Gly Phe Asp Pro Asn Gln Leu Ser
 450 455 460

Val Ala Thr Leu Leu Phe Glu Gly Asp Arg Glu Lys Val Leu Gln His
 465 470 475 480

Glu Lys Gln Val Tyr Asp Ile Ala Ala Lys Phe Gly Gly Leu Ala Ala

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<210> 164
<211> 482
<212> PRT
<213> Homo sapiens

<400> 164
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Met Pro Pro Ser Pro Leu Asp Asp Arg Val Val Val Ala Leu Ser Arg
1 5 10 15

Pro Val Arg Pro Gln Asp Leu Asn Leu Cys Leu Asp Ser Ser Tyr Leu
20 25 30

Gly Ser Ala Asn Pro Gly Ser Asn Ser His Pro Pro Val Ile Ala Thr
35 40 45

Thr Val Val Ser Leu Lys Ala Ala Asn Leu Thr Tyr Met Pro Ser Ser
50 55 60

Ser Gly Ser Ala Arg Ser Leu Asn Cys Gly Cys Ser Ser Ala Ser Cys
65 70 75 80

Cys Thr Val Ala Thr Tyr Asp Lys Asp Asn Gln Ala Gln Thr Gln Ala
85 90 95

Ile Ala Ala Gly Thr Thr Thr Thr Ala Ile Gly Thr Ser Thr Thr Cys
100 105 110

Pro Ala Asn Gln Met Val Asn Asn Asn Glu Asn Thr Gly Ser Leu Ser
115 120 125

Pro Ser Ser Gly Val Gly Ser Pro Val Ser Gly Thr Pro Lys Gln Leu
130 135 140

Ala Ser Ile Lys Ile Ile Tyr Pro Asn Asp Leu Ala Lys Lys Met Thr
145 150 155 160

Lys Cys Ser Lys Ser His Leu Pro Ser Gln Gly Pro Val Ile Ile Asp
165 170 175

Cys Arg Pro Phe Met Glu Tyr Asn Lys Ser His Ile Gln Gly Ala Val
180 185 190

His Ile Asn Cys Ala Asp Lys Ile Ser Arg Arg Arg Leu Gln Gln Gly
195 200 205

Lys Ile Thr Val Leu Asp Leu Ile Ser Cys Arg Glu Gly Lys Asp Ser
210 215 220

Phe Lys Arg Ile Phe Ser Lys Glu Ile Ile Val Tyr Asp Glu Asn Thr
225 230 235 240

236

465 470 475 480

Val Val

<210> 165
 <211> 407
 <212> PRT
 <213> Homo sapiens

<400> 165

Met Glu Ser Ala Ile Thr Leu Trp Gln Phe Leu Leu Gln Leu Leu Leu
 1 5 10 15

Asp Gln Lys His Glu His Leu Ile Cys Trp Thr Ser Asn Asp Gly Glu
 20 25 30

Phe Lys Leu Leu Lys Ala Glu Glu Val Ala Lys Leu Trp Gly Leu Arg
 35 40 45

Lys Asn Lys Thr Asn Met Asn Tyr Asp Lys Leu Ser Arg Ala Leu Arg
 50 55 60

Tyr Tyr Tyr Asp Lys Asn Ile Ile Lys Lys Val Ile Gly Gln Lys Phe
 65 70 75 80

Val Tyr Lys Phe Val Ser Phe Pro Glu Ile Leu Lys Met Asp Pro His
 85 90 95

Ala Val Glu Ile Ser Arg Glu Ser Leu Leu Leu Gln Asp Ser Asp Cys
 100 105 110

Lys Val Ser Pro Glu Gly Arg Glu Ala His Lys His Gly Leu Ala Val
 115 120 125

Leu Arg Ser Thr Ser Arg Asn Glu Tyr Ile His Ser Gly Leu Tyr Ser
 130 135 140

Ser Phe Thr Ile Asn Ser Leu Glu Asn Pro Pro Asp Ala Phe Lys Ala
 145 150 155 160

Ile Lys Arg Glu Lys Leu Glu Glu Pro Pro Glu Asp Ser Pro Pro Val
 165 170 175

Glu Glu Val Arg Thr Val Ile Arg Phe Val Thr Asn Lys Thr Asp Lys
 180 185 190

His Val Thr Arg Pro Val Val Ser Leu Pro Ser Thr Ser Glu Ala Ala
 195 200 205

Ala Ala Ser Ala Phe Leu Ala Ser Ser Val Ser Ala Lys Ile Ser Ser
 210 215 220

Leu Met Leu Pro Asn Ala Ala Ser Ile Ser Ser Ala Ser Pro Phe Ser
 225 230 235 240

Ser Arg Ser Pro Ser Leu Ser Pro Lys Ser Pro Leu Pro Ser Glu His
 245 250 255

Arg Ser Leu Phe Leu Glu Ala Ala Cys His Asp Ser Asp Ser Leu Glu
 260 265 270

Pro Leu Asn Leu Ser Ser Gly Ser Lys Thr Lys Ser Pro Ser Leu Pro
 275 280 285

Pro Lys Ala Lys Lys Pro Lys Gly Leu Glu Ile Ser Ala Pro Pro Leu
 290 295 300

Val Leu Ser Gly Thr Asp Ile Gly Ser Ile Ala Leu Asn Ser Pro Ala
 305 310 315 320

Leu Pro Ser Gly Ser Leu Thr Pro Ala Phe Phe Thr Ala Gln Thr Pro
 325 330 335

Asn Gly Leu Leu Leu Thr Pro Ser Pro Leu Leu Ser Ser Ile His Phe
 340 345 350

Trp Ser Ser Leu Ser Pro Val Ala Pro Leu Ser Pro Ala Arg Leu Gln
 355 360 365

Gly Pro Ser Thr Leu Phe Gln Phe Pro Thr Leu Leu Asn Gly His Met
 370 375 380

Pro Val Pro Ile Pro Ser Leu Asp Arg Ala Ala Ser Pro Val Leu Leu
 385 390 395 400

Ser Ser Asn Ser Gln Lys Ser
405

<210> 166
<211> 364
<212> PRT
<213> Homo sapiens

<400> 166

Met Ala Ala Ile Ser Thr Ser Ile Pro Val Ile Ser Gln Pro Gln Phe
1 5 10 15

Thr Ala Met Asn Glu Pro Gln Cys Phe Tyr Asn Glu Ser Ile Ala Phe
20 25 30

Phe Tyr Asn Arg Ser Gly Lys His Leu Ala Thr Glu Trp Asn Thr Val
35 40 45

Ser Lys Leu Val Met Gly Leu Gly Ile Thr Val Cys Ile Phe Ile Met
50 55 60

Leu Ala Asn Leu Leu Val Met Val Ala Ile Tyr Val Asn Arg Arg Phe
65 70 75 80

His Phe Pro Ile Tyr Tyr Leu Met Ala Asn Leu Ala Ala Ala Asp Phe
85 90 95

Phe Ala Gly Leu Ala Tyr Phe Tyr Leu Met Phe Asn Thr Gly Pro Asn
100 105 110

Thr Arg Arg Leu Thr Val Ser Thr Trp Leu Leu Arg Gln Gly Leu Ile
115 120 125

Asp Thr Ser Leu Thr Ala Ser Val Ala Asn Leu Leu Ala Ile Ala Ile
130 135 140

Glu Arg His Ile Thr Val Phe Arg Met Gln Leu His Thr Arg Met Ser
145 150 155 160

Asn Arg Arg Val Val Val Val Ile Val Val Ile Trp Thr Met Ala Ile
165 170 175

Val Met Gly Ala Ile Pro Ser Val Gly Trp Asn Cys Ile Cys Asp Ile
180 185 190

Glu Asn Cys Ser Asn Met Ala Pro Leu Tyr Ser Asp Ser Tyr Leu Val
 195 200 205

Phe Trp Ala Ile Phe Asn Leu Val Thr Phe Val Val Met Val Val Leu
 210 215 220

Tyr Ala His Ile Phe Gly Tyr Val Arg Gln Arg Thr Met Arg Met Ser
 225 230 235 240

Arg His Ser Ser Gly Pro Arg Arg Asn Arg Asp Thr Met Met Ser Leu
 245 250 255

Leu Lys Thr Val Val Ile Val Leu Gly Ala Phe Ile Ile Cys Trp Thr
 260 265 270

Pro Gly Leu Val Leu Leu Leu Leu Asp Val Cys Cys Pro Gln Cys Asp
 275 280 285

Val Leu Ala Tyr Glu Lys Phe Phe Leu Leu Leu Ala Glu Phe Asn Ser
 290 295 300

Ala Met Asn Pro Ile Ile Tyr Ser Tyr Arg Asp Lys Glu Met Ser Ala
 305 310 315 320

Thr Phe Arg Gln Ile Leu Cys Cys Gln Arg Ser Glu Asn Pro Thr Gly
 325 330 335

Pro Thr Glu Gly Ser Asp Arg Ser Ala Ser Ser Leu Asn His Thr Ile
 340 345 350

Leu Ala Gly Val His Ser Asn Asp His Ser Val Val
 355 360

<210> 167
 <211> 759
 <212> PRT
 <213> Homo sapiens

<400> 167

Met Glu Ser Ser Pro Phe Asn Arg Arg Gln Trp Thr Ser Leu Ser Leu
 1 5 10 15

Arg Val Thr Ala Lys Glu Leu Ser Leu Val Asn Lys Asn Lys Ser Ser
 20 25 30

Ala Ile Val Glu Ile Phe Ser Lys Tyr Gln Lys Ala Ala Glu Glu Thr
 35 40 45

Asn Met Glu Lys Lys Arg Ser Asn Thr Glu Asn Leu Ser Gln His Phe
 50 55 60

Arg Lys Gly Thr Leu Thr Val Leu Lys Lys Lys Trp Glu Asn Pro Gly
 65 70 75 80

Leu Gly Ala Glu Ser His Thr Asp Ser Leu Arg Asn Ser Ser Thr Glu
 85 90 95

Ile Arg His Arg Ala Asp His Pro Pro Ala Glu Val Thr Ser His Ala
 100 105 110

Ala Ser Gly Ala Lys Ala Asp Gln Glu Glu Gln Ile His Pro Arg Ser
 115 120 125

Arg Leu Arg Ser Pro Pro Glu Ala Leu Val Gln Gly Arg Tyr Pro His
 130 135 140

Ile Lys Asp Gly Glu Asp Leu Lys Asp His Ser Thr Glu Ser Lys Lys
 145 150 155 160

Met Glu Asn Cys Leu Gly Glu Ser Arg His Glu Val Glu Lys Ser Glu
 165 170 175

Ile Ser Glu Asn Thr Asp Ala Ser Gly Lys Ile Glu Lys Tyr Asn Val
 180 185 190

Pro Leu Asn Arg Leu Lys Met Met Phe Glu Lys Gly Glu Pro Thr Gln
 195 200 205

Thr Lys Ile Leu Arg Ala Gln Ser Arg Ser Ala Ser Gly Arg Lys Ile
 210 215 220

Ser Glu Asn Ser Tyr Ser Leu Asp Asp Leu Glu Ile Gly Pro Gly Gln
 225 230 235 240

Leu Ser Ser Ser Thr Phe Asp Ser Glu Lys Asn Glu Ser Arg Arg Asn

245	250	255
Leu Glu Leu Pro Arg Leu Ser Glu Thr Ser Ile Lys Asp Arg Met Ala		
260	265	270
Lys Tyr Gln Ala Ala Val Ser Lys Gln Ser Ser Ser Thr Asn Tyr Thr		
275	280	285
Asn Glu Leu Lys Ala Ser Gly Gly Glu Ile Lys Ile His Lys Met Glu		
290	295	300
Gln Lys Glu Asn Val Pro Pro Gly Pro Glu Val Cys Ile Thr His Gln		
305	310	315
Glu Gly Glu Lys Ile Ser Ala Asn Glu Asn Ser Leu Ala Val Arg Ser		
325	330	335
Thr Pro Ala Glu Asp Asp Ser Arg Asp Ser Gln Val Lys Ser Glu Val		
340	345	350
Gln Gln Pro Val His Pro Lys Pro Leu Ser Pro Asp Ser Arg Ala Ser		
355	360	365
Ser Leu Ser Glu Ser Ser Pro Pro Lys Ala Met Lys Lys Phe Gln Ala		
370	375	380
Pro Ala Arg Glu Thr Cys Val Glu Cys Gln Lys Thr Val Tyr Pro Met		
385	390	395
Glu Arg Leu Leu Ala Asn Gln Gln Val Phe His Ile Ser Cys Phe Arg		
405	410	415
Cys Ser Tyr Cys Asn Asn Lys Leu Ser Leu Gly Thr Tyr Ala Ser Leu		
420	425	430
His Gly Arg Ile Tyr Cys Lys Pro His Phe Asn Gln Leu Phe Lys Ser		
435	440	445
Lys Gly Asn Tyr Asp Glu Gly Phe Gly His Arg Pro His Lys Asp Leu		
450	455	460
Trp Ala Ser Lys Asn Glu Asn Glu Glu Ile Leu Glu Arg Pro Ala Gln		
465	470	475
		480

Leu Ala Asn Ala Arg Glu Thr Pro His Ser Pro Gly Val Glu Asp Ala
 485 490 495

Pro Ile Ala Lys Val Gly Val Leu Ala Ala Ser Met Glu Ala Lys Ala
 500 505 510

Ser Ser Gln Gln Glu Lys Glu Asp Lys Pro Ala Glu Thr Lys Lys Leu
 515 520 525

Arg Ile Ala Trp Pro Pro Pro Thr Glu Leu Gly Ser Ser Gly Ser Ala
 530 535 540

Leu Glu Glu Gly Ile Lys Met Ser Lys Pro Lys Trp Pro Pro Glu Asp
 545 550 555 560

Glu Ile Ser Lys Pro Glu Val Pro Glu Asp Val Asp Leu Asp Leu Lys
 565 570 575

Lys Leu Arg Arg Ser Ser Ser Leu Lys Glu Arg Ser Arg Pro Phe Thr
 580 585 590

Val Ala Ala Ser Phe Gln Ser Thr Ser Val Lys Ser Pro Lys Thr Val
 595 600 605

Ser Pro Pro Ile Arg Lys Gly Trp Ser Met Ser Glu Gln Ser Glu Glu
 610 615 620

Ser Val Gly Gly Arg Val Ala Glu Arg Lys Gln Val Glu Asn Ala Lys
 625 630 635 640

Ala Ser Lys Lys Asn Gly Asn Val Gly Lys Thr Thr Trp Gln Asn Lys
 645 650 655

Glu Ser Lys Gly Glu Thr Gly Lys Arg Ser Lys Glu Gly His Ser Leu
 660 665 670

Glu Met Glu Asn Glu Asn Leu Val Glu Asn Gly Ala Asp Ser Asp Glu
 675 680 685

Asp Asp Asn Ser Phe Leu Lys Gln Gln Ser Pro Gln Glu Pro Lys Ser
 690 695 700

Leu Asn Trp Ser Ser Phe Val Asp Asn Thr Phe Ala Glu Glu Phe Thr
705 710 715 720

Thr Gln Asn Gln Lys Ser Gln Asp Val Glu Leu Trp Glu Gly Glu Val
725 730 735

Val Lys Glu Leu Ser Val Glu Glu Gln Ile Lys Arg Asn Arg Tyr Tyr
740 745 750

Asp Glu Asp Glu Asp Glu Glu
755

<210> 168
<211> 695
<212> PRT
<213> Homo sapiens

<400> 168

Met Ile Met Lys Ser Asn Phe Asp Glu Thr Tyr Ile Glu Asn Val Val
1 5 10 15

Arg Asn Ile Leu Lys Gly Gln Asp Val Asp Ser Lys Glu Ala Gln Leu
20 25 30

Ile Ser Phe Leu Ala Leu Leu Ser Ser Tyr Val Thr Asp Ser Thr Ile
35 40 45

Ser Val Ser Gln Cys Glu Ile Phe Leu Gly Ile Ile Tyr Thr Ser Thr
50 55 60

Pro Trp Glu Pro Glu Ser Leu Glu Asp Lys Met Gly Thr Tyr Ser Thr
65 70 75 80

Leu Leu Ile Lys Thr Glu Val Ala Glu Tyr Gly Arg Tyr Thr Gly Val
85 90 95

Arg Ile Ile His Pro Leu Ile Ala Leu Tyr Cys Leu Lys Glu Leu Glu
100 105 110

Arg Ser Tyr His Leu Asp Lys Cys Gln Ile Ala Leu Asn Ile Leu Glu
115 120 125

Glu Asn Leu Phe Tyr Asp Ser Gly Ile Gly Arg Asp Lys Phe Gln His

130 135 140
 Asp Val Gln Thr Leu Leu Leu Thr Arg Gln Arg Lys Val Tyr Gly Asp
 145 150 155 160
 Glu Thr Asp Thr Leu Phe Ser Pro Leu Met Glu Ala Leu Gln Asn Lys
 165 170 175
 Asp Ile Glu Lys Val Leu Ser Ala Gly Ser Arg Arg Phe Pro Gln Asn
 180 185 190
 Ala Phe Ile Cys Gln Ala Leu Ala Arg His Phe Tyr Ile Lys Glu Lys
 195 200 205
 Asp Phe Asn Thr Ala Leu Asp Trp Ala Arg Gln Ala Lys Met Lys Ala
 210 215 220
 Pro Lys Asn Ser Tyr Ile Ser Asp Thr Leu Gly Gln Val Tyr Lys Ser
 225 230 235 240
 Glu Ile Lys Trp Trp Leu Asp Gly Asn Lys Asn Cys Arg Ser Ile Thr
 245 250 255
 Val Asn Asp Leu Thr His Leu Leu Glu Ala Ala Glu Lys Ala Ser Arg
 260 265 270
 Ala Phe Lys Glu Ser Gln Arg Gln Thr Asp Ser Lys Asn Tyr Glu Thr
 275 280 285
 Glu Asn Trp Ser Pro Gln Lys Ser Gln Arg Arg Tyr Asp Met Tyr Asn
 290 295 300
 Thr Ala Cys Phe Leu Gly Glu Ile Glu Val Gly Leu Tyr Thr Ile Gln
 305 310 315 320
 Ile Leu Gln Leu Thr Pro Phe Phe His Lys Glu Asn Glu Leu Ser Lys
 325 330 335
 Lys His Met Val Gln Phe Leu Ser Gly Lys Trp Thr Ile Pro Pro Asp
 340 345 350
 Pro Arg Asn Glu Cys Tyr Leu Ala Leu Ser Lys Phe Thr Ser His Leu
 355 360 365

Lys Asn Leu Gln Ser Asp Leu Lys Arg Cys Phe Asp Phe Phe Ile Asp
 370 375 380

Tyr Met Val Leu Leu Lys Met Arg Tyr Thr Gln Lys Glu Ile Ala Glu
 385 390 395 400

Ile Met Leu Ser Lys Lys Val Ser Arg Cys Phe Arg Lys Tyr Thr Glu
 405 410 415

Leu Phe Cys His Leu Asp Pro Cys Leu Leu Gln Ser Lys Glu Ser Gln
 420 425 430

Leu Leu Gln Glu Glu Asn Cys Arg Lys Lys Leu Glu Ala Leu Arg Ala
 435 440 445

Asp Arg Phe Ala Gly Leu Leu Glu Tyr Leu Asn Pro Asn Tyr Lys Asp
 450 455 460

Ala Thr Thr Met Glu Ser Ile Val Asn Glu Tyr Ala Phe Leu Leu Gln
 465 470 475 480

Gln Asn Ser Lys Lys Pro Met Thr Asn Glu Lys Gln Asn Ser Ile Leu
 485 490 495

Ala Asn Ile Ile Leu Ser Cys Leu Lys Pro Asn Ser Lys Leu Ile Gln
 500 505 510

Pro Leu Thr Thr Leu Lys Lys Gln Leu Arg Glu Val Leu Gln Phe Val
 515 520 525

Gly Leu Ser His Gln Tyr Pro Gly Pro Tyr Phe Leu Ala Cys Leu Leu
 530 535 540

Phe Trp Pro Glu Asn Gln Glu Leu Asp Gln Asp Ser Lys Leu Ile Glu
 545 550 555 560

Lys Tyr Val Ser Ser Leu Asn Arg Ser Phe Arg Gly Gln Tyr Lys Arg
 565 570 575

Met Cys Arg Ser Lys Gln Ala Ser Thr Leu Phe Tyr Leu Gly Lys Arg
 580 585 590

Lys Gly Leu Asn Ser Ile Val His Lys Ala Lys Ile Glu Gln Tyr Phe
 595 600 605

Asp Lys Ala Gln Asn Thr Asn Ser Leu Trp His Ser Gly Asp Val Trp
 610 615 620

Lys Lys Asn Glu Val Lys Asp Leu Leu Arg Arg Leu Thr Gly Gln Ala
 625 630 635 640

Glu Gly Lys Leu Ile Ser Val Glu Tyr Gly Thr Glu Glu Lys Ile Lys
 645 650 655

Ile Pro Val Ile Ser Val Tyr Ser Gly Pro Leu Arg Ser Gly Arg Asn
 660 665 670

Ile Glu Arg Val Ser Phe Tyr Leu Gly Phe Ser Ile Glu Gly Pro Leu
 675 680 685

Ala Tyr Asp Ile Glu Val Ile
 690 695

<210> 169
 <211> 746
 <212> PRT
 <213> Homo sapiens

<400> 169

Met Gln Ala Lys Lys Arg Tyr Phe Ile Leu Leu Ser Ala Gly Ser Cys
 1 5 10 15

Leu Ala Leu Leu Phe Tyr Phe Gly Gly Leu Gln Phe Arg Ala Ser Arg
 20 25 30

Ser His Ser Arg Arg Glu Glu His Ser Gly Arg Asn Gly Leu His His
 35 40 45

Pro Ser Pro Asp His Phe Trp Pro Arg Phe Pro Glu Pro Leu Arg Pro
 50 55 60

Phe Val Pro Trp Asp Gln Leu Glu Asn Glu Asp Ser Ser Val His Ile
 65 70 75 80

Ser Pro Arg Gln Lys Arg Asp Ala Asn Ser Ser Ile Tyr Lys Gly Lys

	85		90		95
Lys Cys Arg Met Glu Ser Cys Phe Asp Phe Thr Leu Cys Lys Lys Asn	100		105		110
Gly Phe Lys Val Tyr Val Tyr Pro Gln Gln Lys Gly Glu Lys Ile Ala	115		120		125
Glu Ser Tyr Gln Asn Ile Leu Ala Ala Ile Glu Gly Ser Arg Phe Tyr	130		135		140
Thr Ser Asp Pro Ser Gln Ala Cys Leu Phe Val Leu Ser Leu Asp Thr	145		150		155
Leu Asp Arg Asp Gln Leu Ser Pro Gln Tyr Val His Asn Leu Arg Ser	165		170		175
Lys Val Gln Ser Leu His Leu Trp Asn Asn Gly Arg Asn His Leu Ile	180		185		190
Phe Asn Leu Tyr Ser Gly Thr Trp Pro Asp Tyr Thr Glu Asp Val Gly	195		200		205
Phe Asp Ile Gly Gln Ala Met Leu Ala Lys Ala Ser Ile Ser Thr Glu	210		215		220
Asn Phe Arg Pro Asn Phe Asp Val Ser Ile Pro Leu Phe Ser Lys Asp	225		230		235
His Pro Arg Thr Gly Gly Glu Arg Gly Phe Leu Lys Phe Asn Thr Ile	245		250		255
Pro Pro Leu Arg Lys Tyr Met Leu Val Phe Lys Gly Lys Arg Tyr Leu	260		265		270
Thr Gly Ile Gly Ser Asp Thr Arg Asn Ala Leu Tyr His Val His Asn	275		280		285
Gly Glu Asp Val Val Leu Leu Thr Thr Cys Lys His Gly Lys Asp Trp	290		295		300
Gln Lys His Lys Asp Ser Arg Cys Asp Arg Asp Asn Thr Glu Tyr Glu	305		310		315
					320

Lys Tyr Asp Tyr Arg Glu Met Leu His Asn Ala Thr Phe Cys Leu Val
 325 330 335

Pro Arg Gly Arg Arg Leu Gly Ser Phe Arg Phe Leu Glu Ala Leu Gln
 340 345 350

Ala Ala Cys Val Pro Val Met Leu Ser Asn Gly Trp Glu Leu Pro Phe
 355 360 365

Ser Glu Val Ile Asn Trp Asn Gln Ala Ala Val Ile Gly Asp Glu Arg
 370 375 380

Leu Leu Leu Gln Ile Pro Ser Thr Ile Arg Ser Ile His Gln Asp Lys
 385 390 395 400

Ile Leu Ala Leu Arg Gln Gln Thr Gln Phe Leu Trp Glu Ala Tyr Phe
 405 410 415

Ser Ser Val Glu Lys Ile Val Leu Thr Thr Leu Glu Ile Ile Gln Asp
 420 425 430

Arg Ile Phe Lys His Ile Ser Arg Asn Ser Leu Ile Trp Asn Lys His
 435 440 445

Pro Gly Gly Leu Phe Val Leu Pro Gln Tyr Ser Ser Tyr Leu Gly Asp
 450 455 460

Phe Pro Tyr Tyr Tyr Ala Asn Leu Gly Leu Lys Pro Pro Ser Lys Phe
 465 470 475 480

Thr Ala Val Ile His Ala Val Thr Pro Leu Val Ser Gln Ser Gln Pro
 485 490 495

Val Leu Lys Leu Leu Val Ala Ala Ala Lys Ser Gln Tyr Cys Ala Gln
 500 505 510

Ile Ile Val Leu Trp Asn Cys Asp Lys Pro Leu Pro Ala Lys His Arg
 515 520 525

Trp Pro Ala Thr Ala Val Pro Val Val Val Ile Glu Gly Glu Ser Lys
 530 535 540

Val Met Ser Ser Arg Phe Leu Pro Tyr Asp Asn Ile Ile Thr Asp Ala
545 550 555 560

Val Leu Ser Leu Asp Glu Asp Thr Val Leu Ser Thr Thr Glu Val Asp
565 570 575

Phe Ala Phe Thr Val Trp Gln Ser Phe Pro Glu Arg Ile Val Gly Tyr
580 585 590

Pro Ala Arg Ser His Phe Trp Asp Asn Ser Lys Glu Arg Trp Gly Tyr
595 600 605

Thr Ser Lys Trp Thr Asn Asp Tyr Ser Met Val Leu Thr Gly Ala Ala
610 615 620

Ile Tyr His Lys Tyr Tyr His Tyr Leu Tyr Ser His Tyr Leu Pro Ala
625 630 635 640

Ser Leu Lys Asn Met Val Asp Gln Leu Ala Asn Cys Glu Asp Ile Leu
645 650 655

Met Asn Phe Leu Val Ser Ala Val Thr Lys Leu Pro Pro Ile Lys Val
660 665 670

Thr Gln Lys Lys Gln Tyr Lys Glu Thr Met Met Gly Gln Thr Ser Arg
675 680 685

Ala Ser Arg Trp Ala Asp Pro Asp His Phe Ala Gln Arg Gln Ser Cys
690 695 700

Met Asn Thr Phe Ala Ser Trp Phe Gly Tyr Met Pro Leu Ile His Ser
705 710 715 720

Gln Met Arg Leu Asp Pro Val Leu Phe Lys Asp Gln Val Ser Ile Leu
725 730 735

Arg Lys Lys Tyr Arg Asp Ile Glu Arg Leu
740 745

<210> 170
<211> 1069
<212> PRT
<213> Homo sapiens

<400> 170

Met Leu Arg Met Arg Thr Ala Gly Trp Ala Arg Gly Trp Cys Leu Gly
 1 5 10 15

Cys Cys Leu Leu Leu Pro Leu Ser Phe Ser Leu Ala Ala Ala Lys Gln
 20 25 30

Leu Leu Arg Tyr Arg Leu Ala Glu Gly Pro Ala Asp Val Arg Ile
 35 40 45

Gly Asn Val Ala Ser Asp Leu Gly Ile Val Thr Gly Ser Gly Glu Val
 50 55 60

Thr Phe Ser Leu Glu Ser Gly Ser Glu Tyr Leu Lys Ile Asp Asn Leu
 65 70 75 80

Thr Gly Glu Leu Ser Thr Ser Glu Arg Arg Ile Asp Arg Glu Lys Leu
 85 90 95

Pro Gln Cys Gln Met Ile Phe Asp Glu Asn Glu Cys Phe Leu Asp Phe
 100 105 110

Glu Val Ser Val Ile Gly Pro Ser Gln Ser Trp Val Asp Leu Phe Glu
 115 120 125

Gly Gln Val Ile Val Leu Asp Ile Asn Asp Asn Thr Pro Thr Phe Pro
 130 135 140

Ser Pro Val Leu Thr Leu Thr Val Glu Glu Asn Arg Pro Val Gly Thr
 145 150 155 160

Leu Tyr Leu Leu Pro Thr Ala Thr Asp Arg Asp Phe Gly Arg Asn Gly
 165 170 175

Ile Glu Arg Tyr Glu Leu Leu Gln Glu Pro Gly Gly Gly Gly Ser Gly
 180 185 190

Gly Glu Ser Arg Arg Ala Gly Ala Ala Asp Ser Ala Pro Tyr Pro Gly
 195 200 205

Gly Gly Gly Asn Gly Ala Ser Gly Gly Gly Ser Gly Gly Ser Lys Arg
 210 215 220

Arg Leu Asp Ala Ser Glu Gly Gly Gly Gly Thr Asn Pro Gly Gly Arg
225 230 235 240

Ser Ser Val Phe Glu Leu Gln Val Ala Asp Thr Pro Asp Gly Glu Lys
245 250 255

Gln Pro Gln Leu Ile Val Lys Gly Ala Leu Asp Arg Glu Gln Arg Asp
260 265 270

Ser Tyr Glu Leu Thr Leu Arg Val Arg Asp Gly Gly Asp Pro Pro Arg
275 280 285

Ser Ser Gln Ala Ile Leu Arg Val Leu Ile Thr Asp Val Asn Asp Asn
290 295 300

Ser Pro Arg Phe Glu Lys Ser Val Tyr Glu Ala Asp Leu Ala Glu Asn
305 310 315 320

Ser Ala Pro Gly Thr Pro Ile Leu Gln Leu Arg Ala Ala Asp Leu Asp
325 330 335

Val Gly Val Asn Gly Gln Ile Glu Tyr Val Phe Gly Ala Ala Thr Glu
340 345 350

Ser Val Arg Arg Leu Leu Arg Leu Asp Glu Thr Ser Gly Trp Leu Ser
355 360 365

Val Leu His Arg Ile Asp Arg Glu Glu Val Asn Gln Leu Arg Phe Thr
370 375 380

Val Met Ala Arg Asp Arg Gly Gln Pro Pro Lys Thr Asp Lys Ala Thr
385 390 395 400

Val Val Leu Asn Ile Lys Asp Glu Asn Asp Asn Val Pro Ser Ile Glu
405 410 415

Ile Arg Lys Ile Gly Arg Ile Pro Leu Lys Asp Gly Val Ala Asn Val
420 425 430

Ala Glu Asp Val Leu Val Asp Thr Pro Ile Ala Leu Val Gln Val Ser
435 440 445

Asp Arg Asp Gln Gly Glu Asn Gly Val Val Thr Cys Thr Val Val Gly
 450 455 460

Asp Val Pro Phe Gln Leu Lys Pro Ala Ser Asp Thr Glu Gly Asp Gln
 465 470 475 480

Asn Lys Lys Lys Tyr Phe Leu His Thr Ser Thr Pro Leu Asp Tyr Glu
 485 490 495

Ala Thr Arg Glu Phe Asn Val Val Ile Val Ala Val Asp Ser Gly Ser
 500 505 510

Pro Ser Leu Ser Ser Lys Asn Ser Leu Ile Val Lys Val Gly Asp Thr
 515 520 525

Asn Asp Asn Pro Pro Met Phe Gly Gln Ser Val Val Glu Val Tyr Phe
 530 535 540

Pro Glu Asn Asn Ile Pro Gly Glu Arg Val Ala Thr Val Leu Ala Thr
 545 550 555 560

Asp Ala Asp Ser Gly Lys Asn Ala Glu Ile Ala Tyr Ser Leu Asp Ser
 565 570 575

Ser Val Met Gly Ile Phe Ala Ile Asp Pro Asp Ser Gly Asp Ile Leu
 580 585 590

Val Asn Thr Val Leu Asp Arg Glu Gln Thr Asp Arg Tyr Glu Phe Lys
 595 600 605

Val Asn Ala Lys Asp Lys Gly Ile Pro Val Leu Gln Gly Ser Thr Thr
 610 615 620

Val Ile Val Gln Val Ala Asp Lys Asn Asp Asn Asp Pro Lys Phe Met
 625 630 635 640

Gln Asp Val Phe Thr Phe Tyr Val Lys Glu Asn Leu Gln Pro Asn Ser
 645 650 655

Pro Val Gly Met Val Thr Val Met Asp Ala Asp Lys Gly Arg Asn Ala
 660 665 670

Glu Met Ser Leu Tyr Ile Glu Glu Asn Asn Asn Ile Phe Ser Ile Glu
675 680 685

Asn Asp Thr Gly Thr Ile Tyr Ser Thr Met Ser Phe Asp Arg Glu His
690 695 700

Gln Thr Thr Tyr Thr Phe Arg Val Lys Ala Val Asp Gly Gly Asp Pro
705 710 715 720

Pro Arg Ser Ala Thr Ala Thr Val Ser Leu Phe Val Met Asp Glu Asn
725 730 735

Asp Asn Ala Pro Thr Val Thr Leu Pro Lys Asn Ile Ser Tyr Thr Leu
740 745 750

Leu Pro Pro Ser Ser Asn Val Arg Thr Val Val Ala Thr Val Leu Ala
755 760 765

Thr Asp Ser Asp Asp Gly Ile Asn Ala Asp Leu Asn Tyr Ser Ile Val
770 775 780

Gly Gly Asn Pro Phe Lys Leu Phe Glu Ile Asp Pro Thr Ser Gly Val
785 790 795 800

Val Ser Leu Val Gly Lys Leu Thr Gln Lys His Tyr Gly Leu His Arg
805 810 815

Leu Val Val Gln Val Asn Asp Ser Gly Gln Pro Ser Gln Ser Thr Thr
820 825 830

Thr Val Val His Val Phe Val Asn Glu Ser Val Ser Asn Ala Thr Ala
835 840 845

Ile Asp Ser Gln Ile Ala Arg Ser Leu His Ile Pro Leu Thr Gln Asp
850 855 860

Ile Ala Gly Asp Pro Ser Tyr Glu Ile Ser Lys Gln Arg Leu Ser Ile
865 870 875 880

Val Ile Gly Val Val Ala Gly Ile Met Thr Val Ile Leu Ile Ile Leu
885 890 895

Ile Val Val Met Ala Arg Tyr Cys Arg Ser Lys Asn Lys Asn Gly Tyr

900 905 910
 Glu Ala Gly Lys Lys Asp His Glu Asp Phe Phe Thr Pro Gln Gln His
 915 920 925
 Asp Lys Ser Lys Lys Pro Lys Lys Asp Lys Lys Asn Lys Lys Ser Lys
 930 935 940
 Gln Pro Leu Tyr Ser Ser Ile Val Thr Val Glu Ala Ser Lys Pro Asn
 945 950 955 960
 Gly Gln Arg Tyr Asp Ser Val Asn Glu Lys Leu Ser Asp Ser Pro Ser
 965 970 975
 Met Gly Arg Tyr Arg Ser Val Asn Gly Gly Pro Gly Ser Pro Asp Leu
 980 985 990
 Ala Arg His Tyr Lys Ser Ser Ser Pro Leu Pro Thr Val Gln Leu His
 995 1000 1005
 Pro Gln Ser Pro Thr Ala Gly Lys Lys His Gln Ala Val Gln Asp
 1010 1015 1020
 Leu Pro Pro Ala Asn Thr Phe Val Gly Ala Gly Asp Asn Ile Ser
 1025 1030 1035
 Ile Gly Ser Asp His Cys Ser Glu Tyr Ser Cys Gln Thr Asn Asn
 1040 1045 1050
 Lys Tyr Ser Lys Gln Met Arg Leu His Pro Tyr Ile Thr Val Phe
 1055 1060 1065

Gly

<210> 171
 <211> 437
 <212> PRT
 <213> Homo sapiens

<400> 171

Met Ser Trp Ser Leu His Pro Arg Asn Leu Ile Leu Tyr Phe Tyr Ala
 1 5 10 15

Leu Leu Phe Leu Ser Ser Thr Cys Val Ala Tyr Val Ala Thr Arg Asp
 20 25 30

Asn Cys Cys Ile Leu Asp Glu Arg Phe Gly Ser Tyr Cys Pro Thr Thr
 35 40 45

Cys Gly Ile Ala Asp Phe Leu Ser Thr Tyr Gln Thr Lys Val Asp Lys
 50 55 60

Asp Leu Gln Ser Leu Glu Asp Ile Leu His Gln Val Glu Asn Lys Thr
 65 70 75 80

Ser Glu Val Lys Gln Leu Ile Lys Ala Ile Gln Leu Thr Tyr Asn Pro
 85 90 95

Asp Glu Ser Ser Lys Pro Asn Met Ile Asp Ala Ala Thr Leu Lys Ser
 100 105 110

Arg Lys Met Leu Glu Glu Ile Met Lys Tyr Glu Ala Ser Ile Leu Thr
 115 120 125

His Asp Ser Ser Ile Arg Tyr Leu Gln Glu Ile Tyr Asn Ser Asn Asn
 130 135 140

Gln Lys Ile Val Asn Leu Lys Glu Lys Val Ala Gln Leu Glu Ala Gln
 145 150 155 160

Cys Gln Glu Pro Cys Lys Asp Thr Val Gln Ile His Asp Ile Thr Gly
 165 170 175

Lys Asp Cys Gln Asp Ile Ala Asn Lys Gly Ala Lys Gln Ser Gly Leu
 180 185 190

Tyr Phe Ile Lys Pro Leu Lys Ala Asn Gln Gln Phe Leu Val Tyr Cys
 195 200 205

Glu Ile Asp Gly Ser Gly Asn Gly Trp Thr Val Phe Gln Lys Arg Leu
 210 215 220

Asp Gly Ser Val Asp Phe Lys Lys Asn Trp Ile Gln Tyr Lys Glu Gly
 225 230 235 240

Phe Gly His Leu Ser Pro Thr Gly Thr Thr Glu Phe Trp Leu Gly Asn
 245 250 255

Glu Lys Ile His Leu Ile Ser Thr Gln Ser Ala Ile Pro Tyr Ala Leu
 260 265 270

Arg Val Glu Leu Glu Asp Trp Asn Gly Arg Thr Ser Thr Ala Asp Tyr
 275 280 285

Ala Met Phe Lys Val Gly Pro Glu Ala Asp Lys Tyr Arg Leu Thr Tyr
 290 295 300

Ala Tyr Phe Ala Gly Gly Asp Ala Gly Asp Ala Phe Asp Gly Phe Asp
 305 310 315 320

Phe Gly Asp Asp Pro Ser Asp Lys Phe Phe Thr Ser His Asn Gly Met
 325 330 335

Gln Phe Ser Thr Trp Asp Asn Asp Asn Asp Lys Phe Glu Gly Asn Cys
 340 345 350

Ala Glu Gln Asp Gly Ser Gly Trp Trp Met Asn Lys Cys His Ala Gly
 355 360 365

His Leu Asn Gly Val Tyr Tyr Gln Gly Gly Thr Tyr Ser Lys Ala Ser
 370 375 380

Thr Pro Asn Gly Tyr Asp Asn Gly Ile Ile Trp Ala Thr Trp Lys Thr
 385 390 395 400

Arg Trp Tyr Ser Met Lys Lys Thr Thr Met Lys Ile Ile Pro Phe Asn
 405 410 415

Arg Leu Thr Ile Gly Glu Gly Gln Gln His His Leu Gly Gly Ala Lys
 420 425 430

Gln Ala Gly Asp Val
 435

<210> 172
 <211> 642
 <212> PRT
 <213> Homo sapiens

<400> 172

Met Lys Arg Ser Ser Val Ser Ser Gly Gly Ala Gly Arg Leu Ser Met
 1 5 10 15

Gln Glu Leu Arg Ser Gln Asp Val Asn Lys Gln Gly Leu Tyr Thr Pro
 20 25 30

Gln Thr Lys Glu Lys Pro Thr Phe Gly Lys Leu Ser Ile Asn Lys Pro
 35 40 45

Thr Ser Glu Arg Lys Val Ser Leu Phe Gly Lys Arg Thr Ser Gly His
 50 55 60

Gly Ser Arg Asn Ser Gln Leu Gly Ile Phe Ser Ser Ser Glu Lys Ile
 65 70 75 80

Lys Asp Pro Arg Pro Leu Asn Asp Lys Ala Phe Ile Gln Gln Cys Ile
 85 90 95

Arg Gln Leu Cys Glu Phe Leu Thr Glu Asn Gly Tyr Ala His Asn Val
 100 105 110

Ser Met Lys Ser Leu Gln Ala Pro Ser Val Lys Asp Phe Leu Lys Ile
 115 120 125

Phe Thr Phe Leu Tyr Gly Phe Leu Cys Pro Ser Tyr Glu Leu Pro Asp
 130 135 140

Thr Lys Phe Glu Glu Glu Val Pro Arg Ile Phe Lys Asp Leu Gly Tyr
 145 150 155 160

Pro Phe Ala Leu Ser Lys Ser Ser Met Tyr Thr Val Gly Ala Pro His
 165 170 175

Thr Trp Pro His Ile Val Ala Ala Leu Val Trp Leu Ile Asp Cys Ile
 180 185 190

Lys Ile His Thr Ala Met Lys Glu Ser Ser Pro Leu Phe Asp Asp Gly
 195 200 205

Gln Pro Trp Gly Glu Glu Thr Glu Asp Gly Ile Met His Asn Lys Leu
 210 215 220

Phe Leu Asp Tyr Thr Ile Lys Cys Tyr Glu Ser Phe Met Ser Gly Ala
 225 230 235 240

Asp Ser Phe Asp Glu Met Asn Ala Glu Leu Gln Ser Lys Leu Lys Asp
 245 250 255

Leu Phe Asn Val Asp Ala Phe Lys Leu Glu Ser Leu Glu Ala Lys Asn
 260 265 270

Arg Ala Leu Asn Glu Gln Ile Ala Arg Leu Glu Gln Glu Arg Glu Lys
 275 280 285

Glu Pro Asn Arg Leu Glu Ser Leu Arg Lys Leu Lys Ala Ser Leu Gln
 290 295 300

Gly Asp Val Gln Lys Tyr Gln Ala Tyr Met Ser Asn Leu Glu Ser His
 305 310 315 320

Ser Ala Ile Leu Asp Gln Lys Leu Asn Gly Leu Asn Glu Glu Ile Ala
 325 330 335

Arg Val Glu Leu Glu Cys Glu Thr Ile Lys Gln Glu Asn Thr Arg Leu
 340 345 350

Gln Asn Ile Ile Asp Asn Gln Lys Tyr Ser Val Ala Asp Ile Glu Arg
 355 360 365

Ile Asn His Glu Arg Asn Glu Leu Gln Gln Thr Ile Asn Lys Leu Thr
 370 375 380

Lys Asp Leu Glu Ala Glu Gln Gln Lys Leu Trp Asn Glu Glu Leu Lys
 385 390 395 400

Tyr Ala Arg Gly Lys Glu Ala Ile Glu Thr Gln Leu Ala Glu Tyr His
 405 410 415

Lys Leu Ala Arg Lys Leu Lys Leu Ile Pro Lys Gly Ala Glu Asn Ser
 420 425 430

Lys Gly Tyr Asp Phe Glu Ile Lys Phe Asn Pro Glu Ala Gly Ala Asn
 435 440 445

Cys Leu Val Lys Tyr Arg Ala Gln Val Tyr Val Pro Leu Lys Glu Leu
 450 455 460

Leu Asn Glu Thr Glu Glu Glu Ile Asn Lys Ala Leu Asn Lys Lys Met
 465 470 475 480

Gly Leu Glu Asp Thr Leu Glu Gln Leu Asn Ala Met Ile Thr Glu Ser
 485 490 495

Lys Arg Ser Val Arg Thr Leu Lys Glu Glu Val Gln Lys Leu Asp Asp
 500 505 510

Leu Tyr Gln Gln Lys Ile Lys Glu Ala Glu Glu Glu Asp Glu Lys Cys
 515 520 525

Ala Ser Glu Leu Glu Ser Leu Glu Lys His Lys His Leu Leu Glu Ser
 530 535 540

Thr Val Asn Gln Gly Leu Ser Glu Ala Met Asn Glu Leu Asp Ala Val
 545 550 555 560

Gln Arg Glu Tyr Gln Leu Val Val Gln Thr Thr Thr Glu Glu Arg Arg
 565 570 575

Lys Val Gly Asn Asn Leu Gln Arg Leu Leu Glu Met Val Ala Thr His
 580 585 590

Val Gly Ser Val Glu Lys His Leu Glu Glu Gln Ile Ala Lys Val Asp
 595 600 605

Arg Glu Tyr Glu Glu Cys Met Ser Glu Asp Leu Ser Glu Asn Ile Lys
 610 615 620

Glu Ile Arg Asp Lys Tyr Glu Lys Lys Ala Thr Leu Ile Lys Ser Ser
 625 630 635 640

Glu Glu

<210> 173
 <211> 178
 <212> PRT
 <213> Homo sapiens

<400> 173

Met Ser Pro Ser Gly Arg Leu Cys Leu Leu Thr Ile Val Gly Leu Ile
 1 5 10 15

Leu Pro Thr Arg Gly Gln Thr Leu Lys Asp Thr Thr Ser Ser Ser Ser
 20 25 30

Ala Asp Ser Thr Ile Met Asp Ile Gln Val Pro Thr Arg Ala Pro Asp
 35 40 45

Ala Val Tyr Thr Glu Leu Gln Pro Thr Ser Pro Thr Pro Thr Trp Pro
 50 55 60

Ala Asp Glu Thr Pro Gln Pro Gln Thr Gln Thr Gln Gln Leu Glu Gly
 65 70 75 80

Thr Asp Gly Pro Leu Val Thr Asp Pro Glu Thr His Lys Ser Thr Lys
 85 90 95

Ala Ala His Pro Thr Asp Asp Thr Thr Thr Leu Ser Glu Arg Pro Ser
 100 105 110

Pro Ser Thr Asp Val Gln Thr Asp Pro Gln Thr Leu Lys Pro Ser Gly
 115 120 125

Phe His Glu Asp Asp Pro Phe Phe Tyr Asp Glu His Thr Leu Arg Lys
 130 135 140

Arg Gly Leu Leu Val Ala Ala Val Leu Phe Ile Thr Gly Ile Ile Ile
 145 150 155 160

Leu Thr Ser Gly Lys Cys Arg Gln Leu Ser Arg Leu Cys Arg Asn His
 165 170 175

Cys Arg

<210> 174

<211> 237

<212> PRT

<213> Homo sapiens

<400> 174

Met Leu Gly Gly Ser Leu Gly Ser Arg Leu Leu Arg Gly Val Gly Gly
1 5 10 15

Ser His Gly Arg Phe Gly Ala Arg Gly Val Arg Glu Gly Gly Ala Ala
20 25 30

Met Ala Ala Gly Glu Ser Met Ala Gln Arg Met Val Trp Val Asp Leu
35 40 45

Glu Met Thr Gly Leu Asp Ile Glu Lys Asp Gln Ile Ile Glu Met Ala
50 55 60

Cys Leu Ile Thr Asp Ser Asp Leu Asn Ile Leu Ala Glu Gly Pro Asn
65 70 75 80

Leu Ile Ile Lys Gln Pro Asp Glu Leu Leu Asp Ser Met Ser Asp Trp
85 90 95

Cys Lys Glu His His Gly Arg Ser Gly Leu Thr Lys Ala Val Lys Glu
100 105 110

Ser Thr Ile Thr Leu Gln Gln Ala Glu Tyr Glu Phe Leu Ser Phe Val
115 120 125

Arg Gln Gln Thr Pro Pro Gly Leu Cys Pro Leu Ala Gly Asn Ser Val
130 135 140

His Glu Asp Lys Lys Phe Leu Asp Lys Tyr Met Pro Gln Phe Met Lys
145 150 155 160

His Leu His Tyr Arg Ile Ile Asp Val Ser Thr Val Lys Glu Leu Cys
165 170 175

Arg Arg Trp Tyr Pro Glu Glu Tyr Glu Phe Ala Pro Lys Lys Ala Ala
180 185 190

Ser His Arg Ala Leu Asp Asp Ile Ser Glu Ser Ile Lys Glu Leu Gln
195 200 205

Phe Tyr Arg Asn Asn Ile Phe Lys Lys Lys Ile Asp Glu Lys Lys Arg
210 215 220

Lys Ile Ile Glu Asn Gly Glu Asn Glu Lys Thr Val Ser

225

230

235

<210> 175

<211> 390

<212> PRT

<213> Homo sapiens

<400> 175

Met Gly Gln Arg Leu Ser Gly Gly Arg Ser Cys Leu Asp Val Pro Gly
 1 5 10 15

Arg Leu Leu Pro Gln Pro Pro Pro Pro Pro Pro Pro Val Arg Arg Lys
 20 25 30

Leu Ala Leu Leu Phe Ala Met Leu Cys Val Trp Leu Tyr Met Phe Leu
 35 40 45

Tyr Ser Cys Ala Gly Ser Cys Ala Ala Ala Pro Gly Leu Leu Leu Leu
 50 55 60

Gly Ser Gly Ser Arg Ala Ala His Asp Pro Pro Ala Leu Ala Thr Ala
 65 70 75 80

Pro Asp Gly Thr Pro Pro Arg Leu Pro Phe Arg Ala Pro Pro Ala Thr
 85 90 95

Pro Leu Ala Ser Gly Lys Glu Met Ala Glu Gly Ala Ala Ser Pro Glu
 100 105 110

Glu Gln Ser Pro Glu Val Pro Asp Ser Pro Ser Pro Ile Ser Ser Phe
 115 120 125

Phe Ser Gly Ser Gly Ser Lys Gln Leu Pro Gln Ala Ile Ile Ile Gly
 130 135 140

Val Lys Lys Gly Gly Thr Arg Ala Leu Leu Glu Phe Leu Arg Val His
 145 150 155 160

Pro Asp Val Arg Ala Val Gly Ala Glu Pro His Phe Phe Asp Arg Ser
 165 170 175

Tyr Asp Lys Gly Leu Ala Trp Tyr Arg Asp Leu Met Pro Arg Thr Leu
 180 185 190

Asp Gly Gln Ile Thr Met Glu Lys Thr Pro Ser Tyr Phe Val Thr Arg
 195 200 205

Glu Ala Pro Ala Arg Ile Ser Ala Met Ser Lys Asp Thr Lys Leu Ile
 210 215 220

Val Val Val Arg Asp Pro Val Thr Arg Ala Ile Ser Asp Tyr Thr Gln
 225 230 235 240

Thr Leu Ser Lys Arg Pro Asp Ile Pro Thr Phe Glu Ser Leu Thr Phe
 245 250 255

Lys Asn Arg Thr Ala Gly Leu Ile Asp Thr Ser Trp Ser Ala Ile Gln
 260 265 270

Ile Gly Ile Tyr Ala Lys His Leu Glu His Trp Leu Arg His Phe Pro
 275 280 285

Ile Arg Gln Met Leu Phe Val Ser Gly Glu Arg Leu Ile Ser Asp Pro
 290 295 300

Ala Gly Glu Leu Gly Arg Val Gln Asp Phe Leu Gly Leu Lys Arg Ile
 305 310 315 320

Ile Thr Asp Lys His Phe Tyr Phe Asn Lys Thr Lys Gly Phe Pro Cys
 325 330 335

Leu Lys Lys Ala Glu Gly Ser Ser Arg Pro His Cys Leu Gly Lys Thr
 340 345 350

Lys Gly Arg Thr His Pro Glu Ile Asp Arg Glu Val Val Arg Arg Leu
 355 360 365

Arg Glu Phe Tyr Arg Pro Phe Asn Leu Lys Phe Tyr Gln Met Thr Gly
 370 375 380

His Asp Phe Gly Trp Asp
 385 390

<210> 176
 <211> 742
 <212> PRT
 <213> Homo sapiens

<400> 176

Met Asp Lys Phe Trp Trp His Ala Ala Trp Gly Leu Cys Leu Val Pro
 1 5 10 15

Leu Ser Leu Ala Gln Ile Asp Leu Asn Ile Thr Cys Arg Phe Ala Gly
 20 25 30

Val Phe His Val Glu Lys Asn Gly Arg Tyr Ser Ile Ser Arg Thr Glu
 35 40 45

Ala Ala Asp Leu Cys Lys Ala Phe Asn Ser Thr Leu Pro Thr Met Ala
 50 55 60

Gln Met Glu Lys Ala Leu Ser Ile Gly Phe Glu Thr Cys Arg Tyr Gly
 65 70 75 80

Phe Ile Glu Gly His Val Val Ile Pro Arg Ile His Pro Asn Ser Ile
 85 90 95

Cys Ala Ala Asn Asn Thr Gly Val Tyr Ile Leu Thr Tyr Asn Thr Ser
 100 105 110

Gln Tyr Asp Thr Tyr Cys Phe Asn Ala Ser Ala Pro Pro Glu Glu Asp
 115 120 125

Cys Thr Ser Val Thr Asp Leu Pro Asn Ala Phe Asp Gly Pro Ile Thr
 130 135 140

Ile Thr Ile Val Asn Arg Asp Gly Thr Arg Tyr Val Gln Lys Gly Glu
 145 150 155 160

Tyr Arg Thr Asn Pro Glu Asp Ile Tyr Pro Ser Asn Pro Thr Asp Asp
 165 170 175

Asp Val Ser Ser Gly Ser Ser Ser Glu Arg Ser Ser Thr Ser Gly Gly
 180 185 190

Tyr Ile Phe Tyr Thr Phe Ser Thr Val His Pro Ile Pro Asp Glu Asp
 195 200 205

Ser Pro Trp Ile Thr Asp Ser Thr Asp Arg Ile Pro Ala Thr Thr Leu
 210 215 220

Met Ser Thr Ser Ala Thr Ala Thr Glu Thr Ala Thr Lys Arg Gln Glu
225 230 235 240

Ala Trp Asp Trp Phe Ser Trp Leu Phe Leu Pro Ser Glu Ser Lys Asn
245 250 255

His Leu His Thr Thr Thr Gln Met Ala Gly Thr Ser Ser Asn Thr Ile
260 265 270

Ser Ala Gly Trp Glu Pro Asn Glu Glu Asn Glu Asp Glu Arg Asp Arg
275 280 285

His Leu Ser Phe Ser Gly Ser Gly Ile Asp Asp Asp Glu Asp Phe Ile
290 295 300

Ser Ser Thr Ile Ser Thr Thr Pro Arg Ala Phe Asp His Thr Lys Gln
305 310 315 320

Asn Gln Asp Trp Thr Gln Trp Asn Pro Ser His Ser Asn Pro Glu Val
325 330 335

Leu Leu Gln Thr Thr Thr Arg Met Thr Asp Val Asp Arg Asn Gly Thr
340 345 350

Thr Ala Tyr Glu Gly Asn Trp Asn Pro Glu Ala His Pro Pro Leu Ile
355 360 365

His His Glu His His Glu Glu Glu Glu Thr Pro His Ser Thr Ser Thr
370 375 380

Ile Gln Ala Thr Pro Ser Ser Thr Thr Glu Glu Thr Ala Thr Gln Lys
385 390 395 400

Glu Gln Trp Phe Gly Asn Arg Trp His Glu Gly Tyr Arg Gln Thr Pro
405 410 415

Arg Glu Asp Ser His Ser Thr Thr Gly Thr Ala Ala Ala Ser Ala His
420 425 430

Thr Ser His Pro Met Gln Gly Arg Thr Thr Pro Ser Pro Glu Asp Ser
435 440 445

Ser Trp Thr Asp Phe Phe Asn Pro Ile Ser His Pro Met Gly Arg Gly
 450 455 460

His Gln Ala Gly Arg Arg Met Asp Met Asp Ser Ser His Ser Thr Thr
 465 470 475 480

Leu Gln Pro Thr Ala Asn Pro Asn Thr Gly Leu Val Glu Asn Leu Asp
 485 490 495

Arg Thr Gly Pro Leu Ser Met Thr Thr Gln Gln Ser Asn Ser Gln Ser
 500 505 510

Phe Ser Thr Ser His Glu Gly Leu Glu Glu Asp Lys Asp His Pro Thr
 515 520 525

Thr Ser Thr Leu Thr Ser Ser Asn Arg Asn Asp Val Thr Gly Gly Arg
 530 535 540

Arg Asp Pro Asn His Ser Glu Gly Ser Thr Thr Leu Leu Glu Gly Tyr
 545 550 555 560

Thr Ser His Tyr Pro His Thr Lys Glu Ser Arg Thr Phe Ile Pro Val
 565 570 575

Thr Ser Ala Lys Thr Gly Ser Phe Gly Val Thr Ala Val Thr Val Gly
 580 585 590

Asp Ser Asn Ser Asn Val Asn Arg Ser Leu Ser Gly Asp Gln Asp Thr
 595 600 605

Phe His Pro Ser Gly Gly Ser His Thr Thr His Gly Ser Glu Ser Asp
 610 615 620

Gly His Ser His Gly Ser Gln Glu Gly Gly Ala Asn Thr Thr Ser Gly
 625 630 635 640

Pro Ile Arg Thr Pro Gln Ile Pro Glu Trp Leu Ile Ile Leu Ala Ser
 645 650 655

Leu Leu Ala Leu Ala Leu Ile Leu Ala Val Cys Ile Ala Val Asn Ser
 660 665 670

Arg Arg Arg Cys Gly Gln Lys Lys Lys Leu Val Ile Asn Ser Gly Asn
675 680 685

Gly Ala Val Glu Asp Arg Lys Pro Ser Gly Leu Asn Gly Glu Ala Ser
690 695 700

Lys Ser Gln Glu Met Val His Leu Val Asn Lys Glu Ser Ser Glu Thr
705 710 715 720

Pro Asp Gln Phe Met Thr Ala Asp Glu Thr Arg Asn Leu Gln Asn Val
725 730 735

Asp Met Lys Ile Gly Val
740

<210> 177
<211> 251
<212> PRT
<213> Homo sapiens

<400> 177

Met Ala Gly Thr Thr Asp Arg Glu Glu Ala Thr Arg Leu Leu Ala Glu
1 5 10 15

Lys Arg Arg Gln Ala Arg Glu Gln Arg Glu Arg Glu Glu Gln Glu Arg
20 25 30

Arg Leu Gln Ala Glu Arg Asp Lys Arg Met Arg Glu Glu Gln Leu Ala
35 40 45

Arg Glu Ala Glu Ala Arg Ala Glu Arg Glu Ala Glu Ala Arg Arg Arg
50 55 60

Glu Glu Gln Glu Ala Arg Glu Lys Ala Gln Ala Glu Gln Glu Glu Gln
65 70 75 80

Glu Arg Leu Gln Lys Gln Lys Glu Glu Ala Glu Ala Arg Ser Arg Glu
85 90 95

Glu Ala Glu Arg Gln Arg Leu Glu Arg Glu Lys His Phe Gln Gln Gln
100 105 110

Glu Gln Glu Arg Gln Glu Arg Arg Lys Arg Leu Glu Glu Ile Met Lys
115 120 125

Arg Thr Arg Lys Ser Glu Val Ser Glu Thr Lys Lys Gln Asp Ser Lys
 130 135 140

Glu Ala Asn Ala Asn Gly Ser Ser Pro Glu Pro Val Lys Ala Val Glu
 145 150 155 160

Ala Arg Ser Pro Gly Leu Gln Lys Glu Ala Val Gln Lys Glu Glu Pro
 165 170 175

Ile Pro Gln Glu Pro Gln Trp Ser Leu Pro Ser Lys Glu Leu Pro Ala
 180 185 190

Ser Leu Val Asn Gly Leu Gln Pro Leu Pro Ala His Gln Glu Asn Gly
 195 200 205

Phe Ser Thr Asn Gly Pro Ser Gly Asp Lys Ser Leu Ser Arg Thr Pro
 210 215 220

Glu Thr Leu Leu Pro Phe Ala Glu Ala Glu Ala Phe Leu Lys Lys Ala
 225 230 235 240

Val Val Gln Ser Pro Gln Val Thr Glu Val Leu
 245 250

<210> 178
 <211> 71
 <212> PRT
 <213> Homo sapiens

<400> 178

Ser Ser Lys Thr Ala Ser Thr Asn Asn Ile Ala Gln Ala Arg Arg Thr
 1 5 10 15

Val Gln Gln Leu Arg Leu Glu Ala Ser Ile Glu Arg Ile Lys Val Ser
 20 25 30

Lys Ala Ser Ala Asp Leu Met Ser Tyr Cys Glu Glu His Ala Arg Ser
 35 40 45

Asp Pro Leu Leu Ile Gly Ile Pro Thr Ser Glu Asn Pro Phe Lys Asp
 50 55 60

Lys Lys Thr Cys Ile Ile Leu
65 70

<210> 179
<211> 292
<212> PRT
<213> Homo sapiens

<400> 179

Met Asn Leu Asn Met Gly Arg Glu Met Lys Glu Glu Leu Glu Glu Glu
1 5 10 15

Glu Lys Met Arg Glu Asp Gly Gly Gly Lys Asp Arg Ala Lys Ser Lys
20 25 30

Lys Val His Arg Ile Val Ser Lys Trp Met Leu Pro Glu Lys Ser Arg
35 40 45

Gly Thr Tyr Leu Glu Arg Ala Asn Cys Phe Pro Pro Pro Val Phe Ile
50 55 60

Ile Ser Ile Ser Leu Ala Glu Leu Ala Val Phe Ile Tyr Tyr Ala Val
65 70 75 80

Trp Lys Pro Gln Lys Gln Trp Ile Thr Leu Asp Thr Gly Ile Leu Glu
85 90 95

Ser Pro Phe Ile Tyr Ser Pro Glu Lys Arg Glu Glu Ala Trp Arg Phe
100 105 110

Ile Ser Tyr Met Leu Val His Ala Gly Val Gln His Ile Leu Gly Asn
115 120 125

Leu Cys Met Gln Leu Val Leu Gly Ile Pro Leu Glu Met Val His Lys
130 135 140

Gly Leu Arg Val Gly Leu Val Tyr Leu Ala Gly Val Ile Ala Gly Ser
145 150 155 160

Leu Ala Ser Ser Ile Phe Asp Pro Leu Arg Tyr Leu Val Gly Ala Ser
165 170 175

Gly Gly Val Tyr Ala Leu Met Gly Gly Tyr Phe Met Asn Val Leu Val
180 185 190

Asn Phe Gln Glu Met Ile Pro Ala Phe Gly Ile Phe Arg Leu Leu Ile
 195 200 205

Ile Ile Leu Ile Ile Val Leu Asp Met Gly Phe Ala Leu Tyr Arg Arg
 210 215 220

Phe Phe Val Pro Glu Asp Gly Ser Pro Val Ser Phe Ala Ala His Ile
 225 230 235 240

Ala Gly Gly Phe Ala Gly Met Ser Ile Gly Tyr Thr Val Phe Ser Cys
 245 250 255

Phe Asp Lys Ala Leu Leu Lys Asp Pro Arg Phe Trp Ile Ala Ile Ala
 260 265 270

Ala Tyr Leu Ala Cys Val Leu Phe Ala Val Phe Phe Asn Ile Phe Leu
 275 280 285

Ser Pro Ala Asn
 290

<210> 180
 <211> 775
 <212> PRT
 <213> Homo sapiens

<400> 180

Met Ala Ser Arg Ala Val Val Arg Ala Arg Arg Cys Pro Gln Cys Pro
 1 5 10 15

Gln Val Arg Ala Ala Ala Ala Ala Pro Ala Trp Ala Ala Leu Pro Leu
 20 25 30

Ser Arg Ser Leu Pro Pro Cys Ser Asn Ser Ser Ser Phe Ser Met Pro
 35 40 45

Leu Phe Leu Leu Leu Leu Val Leu Leu Leu Leu Leu Glu Asp Ala
 50 55 60

Gly Ala Gln Gln Gly Asp Gly Cys Gly His Thr Val Leu Gly Pro Glu
 65 70 75 80

Ser Gly Thr Leu Thr Ser Ile Asn Tyr Pro Gln Thr Tyr Pro Asn Ser
85 90 95

Thr Val Cys Glu Trp Glu Ile Arg Val Lys Met Gly Glu Arg Val Arg
100 105 110

Ile Lys Phe Gly Asp Phe Asp Ile Glu Asp Ser Asp Ser Cys His Phe
115 120 125

Asn Tyr Leu Arg Ile Tyr Asn Gly Ile Gly Val Ser Arg Thr Glu Ile
130 135 140

Gly Lys Tyr Cys Gly Leu Gly Leu Gln Met Asn His Ser Ile Glu Ser
145 150 155 160

Lys Gly Asn Glu Ile Thr Leu Leu Phe Met Ser Gly Ile His Val Ser
165 170 175

Gly Arg Gly Phe Leu Ala Ser Tyr Ser Val Ile Asp Lys Gln Asp Leu
180 185 190

Ile Thr Cys Leu Asp Thr Ala Ser Asn Phe Leu Glu Pro Glu Phe Ser
195 200 205

Lys Tyr Cys Pro Ala Gly Cys Leu Leu Pro Phe Ala Glu Ile Ser Gly
210 215 220

Thr Ile Pro His Gly Tyr Arg Asp Ser Ser Pro Leu Cys Met Ala Gly
225 230 235 240

Val His Ala Gly Val Val Ser Asn Thr Leu Gly Gly Gln Ile Ser Val
245 250 255

Val Ile Ser Lys Gly Ile Pro Tyr Tyr Glu Ser Ser Leu Ala Asn Asn
260 265 270

Val Thr Ser Val Val Gly His Leu Ser Thr Ser Leu Phe Thr Phe Lys
275 280 285

Thr Ser Gly Cys Tyr Gly Thr Leu Gly Met Glu Ser Gly Val Ile Ala
290 295 300

Asp Pro Gln Ile Thr Ala Ser Ser Val Leu Glu Trp Thr Asp His Thr

305 310 315 320
 Gly Gln Glu Asn Ser Trp Lys Pro Lys Lys Ala Arg Leu Lys Lys Pro
 325 330 335
 Gly Pro Pro Trp Ala Ala Phe Ala Thr Asp Glu Tyr Gln Trp Leu Gln
 340 345 350
 Ile Asp Leu Asn Lys Glu Lys Lys Ile Thr Gly Ile Ile Thr Thr Gly
 355 360 365
 Ser Thr Met Val Glu His Asn Tyr Tyr Val Ser Ala Tyr Arg Ile Leu
 370 375 380
 Tyr Ser Asp Asp Gly Gln Lys Trp Thr Val Tyr Arg Glu Pro Gly Val
 385 390 395 400
 Glu Gln Asp Lys Ile Phe Gln Gly Asn Lys Asp Tyr His Gln Asp Val
 405 410 415
 Arg Asn Asn Phe Leu Pro Pro Ile Ile Ala Arg Phe Ile Arg Val Asn
 420 425 430
 Pro Thr Gln Trp Gln Gln Lys Ile Ala Met Lys Met Glu Leu Leu Gly
 435 440 445
 Cys Gln Phe Ile Pro Lys Gly Arg Pro Pro Lys Leu Thr Gln Pro Pro
 450 455 460
 Pro Pro Arg Asn Ser Asn Asp Leu Lys Asn Thr Thr Ala Pro Pro Lys
 465 470 475 480
 Ile Ala Lys Gly Arg Ala Pro Lys Phe Thr Gln Pro Leu Gln Pro Arg
 485 490 495
 Ser Ser Asn Glu Phe Pro Ala Gln Thr Glu Gln Thr Thr Ala Ser Pro
 500 505 510
 Asp Ile Arg Asn Thr Thr Val Thr Pro Asn Val Thr Lys Asp Val Ala
 515 520 525
 Leu Ala Ala Val Leu Val Pro Val Leu Val Met Val Leu Thr Thr Leu
 530 535 540

Ile Leu Ile Leu Val Cys Ala Trp His Trp Arg Asn Arg Lys Lys Lys
 545 550 555 560

Thr Glu Gly Thr Tyr Asp Leu Pro Tyr Trp Asp Arg Ala Gly Trp Trp
 565 570 575

Lys Gly Met Lys Gln Phe Leu Pro Ala Lys Ala Val Asp His Glu Glu
 580 585 590

Thr Pro Val Arg Tyr Ser Ser Ser Glu Val Asn His Leu Ser Pro Arg
 595 600 605

Glu Val Thr Thr Val Leu Gln Ala Asp Ser Ala Glu Tyr Ala Gln Pro
 610 615 620

Leu Val Gly Gly Ile Val Gly Thr Leu His Gln Arg Ser Thr Phe Lys
 625 630 635 640

Pro Glu Glu Gly Lys Glu Ala Gly Tyr Ala Asp Leu Asp Pro Tyr Asn
 645 650 655

Ser Pro Gly Gln Glu Val Tyr His Ala Tyr Ala Glu Pro Leu Pro Ile
 660 665 670

Thr Gly Pro Glu Tyr Ala Thr Pro Ile Ile Met Asp Met Ser Gly His
 675 680 685

Pro Thr Thr Ser Val Gly Gln Pro Ser Thr Ser Thr Phe Lys Ala Thr
 690 695 700

Gly Asn Gln Pro Pro Pro Leu Val Gly Thr Tyr Asn Thr Leu Leu Ser
 705 710 715 720

Arg Thr Asp Ser Cys Ser Ser Ala Gln Ala Gln Tyr Asp Thr Pro Lys
 725 730 735

Ala Gly Lys Pro Gly Leu Pro Ala Pro Asp Glu Leu Val Tyr Gln Val
 740 745 750

Pro Gln Ser Thr Gln Glu Val Ser Gly Ala Gly Arg Asp Gly Glu Cys
 755 760 765

Asp Val Phe Lys Glu Ile Leu
770 775

<210> 181
<211> 494
<212> PRT
<213> Homo sapiens

<400> 181

Glu Asn Tyr Lys Asn Leu Val Ala Val Asp Trp Glu Ser His Ile Asn
1 5 10 15

Thr Lys Trp Ser Ala Pro Gln Gln Asn Phe Leu Gln Gly Lys Thr Ser
20 25 30

Ser Val Val Glu Met Glu Arg Asn His Phe Gly Glu Glu Leu Phe Asp
35 40 45

Phe Asn Gln Cys Glu Lys Ala Leu Ser Glu His Ser Cys Leu Lys Thr
50 55 60

His Arg Arg Thr Tyr Phe Arg Lys Lys Thr Cys Glu Cys Asn Gln Cys
65 70 75 80

Glu Lys Ala Phe Arg Lys Pro Ser Ile Phe Thr Leu His Lys Lys Thr
85 90 95

Asp Ile Gly Glu Glu Leu Pro Asn Cys Asn Gln Cys Glu Thr Ala Phe
100 105 110

Ser Gln His Leu His Leu Val Cys Lys Lys Thr Ser Gln Asn Leu His
115 120 125

Leu Val Cys Lys Lys Thr His Thr Gln Glu Lys Pro Tyr Lys Cys Ser
130 135 140

Asp Cys Glu Lys Gly Leu Pro Ser Ser Ser His Leu Arg Glu Cys Val
145 150 155 160

Arg Ile Tyr Gly Gly Glu Arg Pro Tyr Thr His Lys Glu Tyr Val Glu
165 170 175

Thr Phe Ser His Ser Thr Ala Leu Phe Val His Met Gln Thr Gln Asp

180	185	190
Gly Glu Lys Phe Tyr Glu Cys Lys Ala Cys Gly Lys Pro Phe Thr Glu		
195	200	205
Ser Ser Tyr Leu Thr Gln His Leu Arg Thr His Ser Arg Val Leu Pro		
210	215	220
Ile Glu His Lys Lys Phe Gly Lys Ala Phe Ala Phe Ser Pro Asp Leu		
225	230	235 240
Ala Lys His Ile Arg Leu Arg Thr Arg Gly Lys His Tyr Val Cys Asn		
245	250	255
Glu Cys Gly Lys Glu Phe Thr Cys Phe Ser Lys Leu Asn Ile His Ile		
260	265	270
Arg Val His Thr Gly Glu Lys Pro Tyr Glu Cys Asn Lys Cys Gly Lys		
275	280	285
Ala Phe Thr Asp Ser Ser Gly Leu Ile Lys His Arg Arg Thr His Thr		
290	295	300
Gly Glu Lys Pro Tyr Glu Cys Lys Glu Cys Gly Lys Ala Phe Ala Asn		
305	310	315 320
Ser Ser His Leu Thr Val His Met Arg Thr His Thr Gly Glu Lys Pro		
325	330	335
Tyr Gln Cys Lys Glu Cys Gly Lys Ala Phe Ile Asn Ser Ser Ser Phe		
340	345	350
Lys Ser His Met Gln Thr His Pro Gly Val Lys Pro Tyr Asp Cys Gln		
355	360	365
Gln Cys Gly Lys Ala Phe Ile Arg Ser Ser Phe Leu Ile Arg His Leu		
370	375	380
Arg Ser His Ser Ala Glu Arg Pro Phe Glu Cys Glu Glu Cys Gly Lys		
385	390	395 400
Ala Phe Arg Tyr Ser Ser His Leu Ser Gln His Lys Arg Ile His Thr		
405	410	415

Gly Glu Arg Pro Tyr Lys Cys Gln Lys Cys Gly Gln Ala Phe Ser Ile
 420 425 430

Ser Ser Gly Leu Thr Val His Met Arg Thr His Thr Gly Glu Arg Pro
 435 440 445

Phe Glu Cys Gln Glu Cys Gly Lys Ala Phe Thr Arg Ser Thr Tyr Leu
 450 455 460

Ile Arg His Leu Arg Ser His Ser Val Glu Lys Pro Tyr Lys Glu Cys
 465 470 475 480

Gly Gln Thr Phe Ser Asn Ser Ser Cys Leu Thr Glu Cys Val
 485 490

<210> 182
 <211> 556
 <212> PRT
 <213> Homo sapiens

<400> 182

Met Met Asn Lys Leu Tyr Ile Gly Asn Leu Ser Pro Ala Val Thr Ala
 1 5 10 15

Asp Asp Leu Arg Gln Leu Phe Gly Asp Arg Lys Leu Pro Leu Ala Gly
 20 25 30

Gln Val Leu Leu Lys Ser Gly Tyr Ala Phe Val Asp Tyr Pro Asp Gln
 35 40 45

Asn Trp Ala Ile Arg Ala Ile Glu Thr Leu Ser Gly Lys Val Glu Leu
 50 55 60

His Gly Lys Ile Met Glu Val Asp Tyr Ser Val Ser Lys Lys Leu Arg
 65 70 75 80

Ser Arg Lys Ile Gln Ile Arg Asn Ile Pro Pro His Leu Gln Trp Glu
 85 90 95

Val Leu Asp Gly Leu Leu Ala Gln Tyr Gly Thr Val Glu Asn Val Glu
 100 105 110

Gln Val Asn Thr Asp Thr Glu Thr Ala Val Val Asn Val Thr Tyr Ala
 115 120 125

Thr Arg Glu Glu Ala Lys Ile Ala Met Glu Lys Leu Ser Gly His Gln
 130 135 140

Phe Glu Asn Tyr Ser Phe Lys Ile Ser Tyr Ile Pro Asp Glu Glu Val
 145 150 155 160

Ser Ser Pro Ser Pro Pro Gln Arg Ala Gln Arg Gly Asp His Ser Ser
 165 170 175

Arg Glu Gln Gly His Ala Pro Gly Gly Thr Ser Gln Ala Arg Gln Ile
 180 185 190

Asp Phe Pro Leu Arg Ile Leu Val Pro Thr Gln Phe Val Gly Ala Ile
 195 200 205

Ile Gly Lys Glu Gly Leu Thr Ile Lys Asn Ile Thr Lys Gln Thr Gln
 210 215 220

Ser Arg Val Asp Ile His Arg Lys Glu Asn Ser Gly Ala Ala Glu Lys
 225 230 235 240

Pro Val Thr Ile His Ala Thr Pro Glu Gly Thr Ser Glu Ala Cys Arg
 245 250 255

Met Ile Leu Glu Ile Met Gln Lys Glu Ala Asp Glu Thr Lys Leu Ala
 260 265 270

Glu Glu Ile Pro Leu Lys Ile Leu Ala His Asn Gly Leu Val Gly Arg
 275 280 285

Leu Ile Gly Lys Glu Gly Arg Asn Leu Lys Lys Ile Glu His Glu Thr
 290 295 300

Gly Thr Lys Ile Thr Ile Ser Ser Leu Gln Asp Leu Ser Ile Tyr Asn
 305 310 315 320

Pro Glu Arg Thr Ile Thr Val Lys Gly Thr Val Glu Ala Cys Ala Ser
 325 330 335

Ala Glu Ile Glu Ile Met Lys Lys Leu Arg Glu Ala Phe Glu Asn Asp

340	345	350
Met Leu Ala Val Asn Thr His Ser Gly Tyr Phe Ser Ser Leu Tyr Pro		
355	360	365
His His Gln Phe Gly Pro Phe Pro His His His Ser Tyr Pro Glu Gln		
370	375	380
Glu Ile Val Asn Leu Phe Ile Pro Thr Gln Ala Val Gly Ala Ile Ile		
385	390	395
Gly Lys Lys Gly Ala His Ile Lys Gln Leu Ala Arg Phe Ala Gly Ala		
405	410	415
Ser Ile Lys Ile Ala Pro Ala Glu Gly Pro Asp Val Ser Glu Arg Met		
420	425	430
Val Ile Ile Thr Gly Pro Pro Glu Ala Gln Phe Lys Ala Gln Gly Arg		
435	440	445
Ile Phe Gly Lys Leu Lys Glu Glu Asn Phe Phe Asn Pro Lys Glu Glu		
450	455	460
Val Lys Leu Glu Ala His Ile Arg Val Pro Ser Ser Thr Ala Gly Arg		
465	470	475
Val Ile Gly Lys Gly Gly Lys Thr Val Asn Glu Leu Gln Asn Leu Thr		
485	490	495
Ser Ala Glu Val Ile Val Pro Arg Asp Gln Thr Pro Asp Glu Asn Glu		
500	505	510
Glu Val Ile Val Arg Ile Ile Gly His Phe Phe Ala Ser Gln Thr Ala		
515	520	525
Gln Arg Lys Ile Arg Glu Ile Val Gln Gln Val Lys Gln Gln Glu Gln		
530	535	540
Lys Tyr Pro Gln Gly Val Ala Ser Gln Arg Ser Lys		
545	550	555
<210> 183		
<211> 399		

<212> PRT

<213> Homo sapiens

<400> 183

Met Ser Asp Ile Leu Arg Glu Leu Leu Cys Val Ser Glu Lys Ala Ala
 1 5 10 15

Asn Ile Ala Arg Ala Cys Arg Gln Gln Glu Ala Leu Phe Gln Leu Leu
 20 25 30

Ile Glu Glu Lys Lys Glu Gly Glu Lys Asn Lys Lys Phe Ala Val Asp
 35 40 45

Phe Lys Thr Leu Ala Asp Val Leu Val Gln Glu Val Ile Lys Gln Asn
 50 55 60

Met Glu Asn Lys Phe Pro Gly Leu Glu Lys Asn Ile Phe Gly Glu Glu
 65 70 75 80

Ser Asn Glu Phe Thr Asn Asp Trp Gly Glu Lys Ile Thr Leu Arg Leu
 85 90 95

Cys Ser Thr Glu Glu Glu Thr Ala Glu Leu Leu Ser Lys Val Leu Asn
 100 105 110

Gly Asn Lys Val Ala Ser Glu Ala Leu Ala Arg Val Val His Gln Asp
 115 120 125

Val Ala Phe Thr Asp Pro Thr Leu Asp Ser Thr Glu Ile Asn Val Pro
 130 135 140

Gln Asp Ile Leu Gly Ile Trp Val Asp Pro Ile Asp Ser Thr Tyr Gln
 145 150 155 160

Tyr Ile Lys Gly Ser Ala Asp Ile Lys Ser Asn Gln Gly Ile Phe Pro
 165 170 175

Cys Gly Leu Gln Cys Val Thr Ile Leu Ile Gly Val Tyr Asp Ile Gln
 180 185 190

Thr Gly Val Pro Leu Met Gly Val Ile Asn Gln Pro Phe Val Ser Arg
 195 200 205

Asp Pro Asn Thr Leu Arg Trp Lys Gly Gln Cys Tyr Trp Gly Leu Ser
 210 215 220

Tyr Met Gly Thr Asn Met His Ser Leu Gln Leu Thr Ile Ser Arg Arg
 225 230 235 240

Asn Gly Ser Glu Thr His Thr Gly Asn Thr Gly Ser Glu Ala Ala Phe
 245 250 255

Ser Pro Ser Phe Ser Ala Val Ile Ser Thr Ser Glu Lys Glu Thr Ile
 260 265 270

Lys Ala Ala Leu Ser Arg Val Cys Gly Asp Arg Ile Phe Gly Ala Ala
 275 280 285

Gly Ala Gly Tyr Lys Ser Leu Cys Val Val Gln Gly Leu Val Asp Ile
 290 295 300

Tyr Ile Phe Ser Glu Asp Thr Thr Phe Lys Trp Asp Ser Cys Ala Ala
 305 310 315 320

His Ala Ile Leu Arg Ala Met Gly Gly Gly Ile Val Asp Leu Lys Glu
 325 330 335

Cys Leu Glu Arg Asn Pro Glu Thr Gly Leu Asp Leu Pro Gln Leu Val
 340 345 350

Tyr His Val Glu Asn Glu Gly Ala Ala Gly Val Asp Arg Trp Ala Asn
 355 360 365

Lys Gly Gly Leu Ile Ala Tyr Arg Ser Arg Lys Arg Leu Glu Thr Phe
 370 375 380

Leu Ser Leu Leu Val Gln Asn Leu Ala Pro Ala Glu Thr His Thr
 385 390 395

<210> 184
 <211> 662
 <212> PRT
 <213> Homo sapiens

<400> 184

Pro Leu Cys Pro Ala Leu Cys Pro Thr Ser Pro Pro Pro Leu Pro Leu
 1 5 10 15

Leu Pro Pro Ser Val Ser Pro Pro Gly Cys Leu Thr Leu Trp Ser Leu
 20 25 30

Ser Phe Leu Phe Ser Val Pro Ser Ala Pro Tyr Pro His Leu Lys Thr
 35 40 45

Thr Met Ala Thr Ile Pro Asp Trp Lys Leu Gln Leu Leu Ala Arg Arg
 50 55 60

Arg Gln Glu Glu Ala Ser Val Arg Gly Arg Glu Lys Ala Glu Arg Glu
 65 70 75 80

Arg Leu Ser Gln Met Pro Ala Trp Lys Arg Gly Leu Leu Glu Arg Arg
 85 90 95

Arg Ala Lys Leu Gly Leu Ser Pro Gly Glu Pro Ser Pro Val Leu Gly
 100 105 110

Thr Val Glu Ala Gly Pro Pro Asp Pro Asp Glu Ser Ala Val Leu Leu
 115 120 125

Glu Ala Ile Gly Pro Val His Gln Asn Arg Phe Ile Arg Gln Glu Arg
 130 135 140

Gln Gln Gln Gln Gln Gln Gln Gln Arg Ser Glu Glu Leu Leu Ala Glu
 145 150 155 160

Arg Lys Pro Gly Pro Leu Glu Ala Arg Glu Arg Arg Pro Ser Pro Gly
 165 170 175

Glu Met Arg Asp Gln Ser Pro Lys Gly Arg Glu Ser Arg Glu Glu Arg
 180 185 190

Leu Ser Pro Arg Glu Thr Arg Glu Arg Arg Leu Gly Ile Gly Gly Ala
 195 200 205

Gln Glu Leu Ser Leu Arg Pro Leu Glu Ala Arg Asp Trp Arg Gln Ser
 210 215 220

Pro Gly Glu Val Gly Asp Arg Ser Ser Arg Leu Ser Glu Ala Trp Lys
 225 230 235 240

Pro Ser Pro Leu Pro Pro Glu Asp Ala Gly Thr Gly Gly Leu Arg Gln
450 455 460

Gln Glu Glu Glu Ala Val Glu Leu Gln Pro Pro Pro Pro Ala Pro Leu
 465 470 475 480

Ser Pro Pro Pro Pro Ala Pro Thr Ala Pro Gln Pro Pro Gly Asp Pro
 485 490 495

Leu Met Ser Arg Leu Phe Tyr Gly Val Lys Ala Gly Pro Gly Val Gly
 500 505 510

Ala Pro Arg Arg Ser Gly His Thr Phe Thr Val Asn Pro Arg Arg Ser
 515 520 525

Val Pro Pro Ala Thr Pro Ala Thr Pro Thr Ser Pro Ala Thr Val Asp
 530 535 540

Ala Ala Val Pro Gly Ala Gly Lys Lys Arg Tyr Pro Thr Ala Glu Glu
 545 550 555 560

Ile Leu Val Leu Gly Gly Tyr Leu Arg Leu Ser Arg Ser Cys Leu Ala
 565 570 575

Lys Gly Ser Pro Glu Arg His His Lys Gln Leu Lys Ile Ser Phe Ser
 580 585 590

Glu Thr Ala Leu Glu Thr Thr Tyr Gln Tyr Pro Ser Glu Ser Ser Val
 595 600 605

Leu Glu Glu Leu Gly Pro Glu Pro Glu Val Pro Ser Ala Pro Asn Pro
 610 615 620

Pro Ala Ala Gln Pro Asp Asp Glu Glu Asp Glu Glu Glu Leu Leu Leu
 625 630 635 640

Leu Gln Pro Glu Leu Gln Gly Gly Leu Arg Thr Lys Ala Leu Ile Val
 645 650 655

Asp Glu Ser Cys Arg Arg
 660

<210> 185
 <211> 1609
 <212> PRT
 <213> Homo sapiens

<400> 185

Met Arg Gly Ser His Arg Ala Ala Pro Ala Leu Arg Pro Arg Gly Arg
 1 5 10 15

Leu Trp Pro Val Leu Ala Val Leu Ala Ala Ala Ala Ala Gly Cys
 20 25 30

Ala Gln Ala Ala Met Asp Glu Cys Thr Asp Glu Gly Gly Arg Pro Gln
 35 40 45

Arg Cys Met Pro Glu Phe Val Asn Ala Ala Phe Asn Val Thr Val Val
 50 55 60

Ala Thr Asn Thr Cys Gly Thr Pro Pro Glu Glu Tyr Cys Val Gln Thr
 65 70 75 80

Gly Val Thr Gly Val Thr Lys Ser Cys His Leu Cys Asp Ala Gly Gln
 85 90 95

Pro His Leu Gln His Gly Ala Ala Phe Leu Thr Asp Tyr Asn Asn Gln
 100 105 110

Ala Asp Thr Thr Trp Trp Gln Ser Gln Thr Met Leu Ala Gly Val Gln
 115 120 125

Tyr Pro Ser Ser Ile Asn Leu Thr Leu His Leu Gly Lys Ala Phe Asp
 130 135 140

Ile Thr Tyr Val Arg Leu Lys Phe His Thr Ser Arg Pro Glu Ser Phe
 145 150 155 160

Ala Ile Tyr Lys Arg Thr Arg Glu Asp Gly Pro Trp Ile Pro Tyr Gln
 165 170 175

Tyr Tyr Ser Gly Ser Cys Glu Asn Thr Tyr Ser Lys Ala Asn Arg Gly
 180 185 190

Phe Ile Arg Thr Gly Gly Asp Glu Gln Gln Ala Leu Cys Thr Asp Glu
 195 200 205

Phe Ser Asp Phe Ser Pro Leu Thr Gly Gly Asn Val Ala Phe Ser Thr
 210 215 220

Leu Glu Gly Arg Pro Ser Ala Tyr Asn Phe Asp Asn Ser Pro Val Leu
 225 230 235 240

Gln Glu Trp Val Thr Ala Thr Asp Ile Arg Val Thr Leu Asn Arg Leu
 245 250 255

Asn Thr Phe Gly Asp Glu Val Phe Asn Asp Pro Lys Val Leu Lys Ser
 260 265 270

Tyr Tyr Tyr Ala Ile Ser Asp Phe Ala Val Gly Gly Arg Cys Lys Cys
 275 280 285

Asn Gly His Ala Ser Glu Cys Met Lys Asn Glu Phe Asp Lys Leu Val
 290 295 300

Cys Asn Cys Lys His Asn Thr Tyr Gly Val Asp Cys Glu Lys Cys Leu
 305 310 315 320

Pro Phe Phe Asn Asp Arg Pro Trp Arg Arg Ala Thr Ala Glu Ser Ala
 325 330 335

Ser Glu Cys Leu Pro Cys Asp Cys Asn Gly Arg Ser Gln Glu Cys Tyr
 340 345 350

Phe Asp Pro Glu Leu Tyr Arg Ser Thr Gly His Gly Gly His Cys Thr
 355 360 365

Asn Cys Gln Asp Asn Thr Asp Gly Ala His Cys Glu Arg Cys Arg Glu
 370 375 380

Asn Phe Phe Arg Leu Gly Asn Asn Glu Ala Cys Ser Ser Cys His Cys
 385 390 395 400

Ser Pro Val Gly Ser Leu Ser Thr Gln Cys Asp Ser Tyr Gly Arg Cys
 405 410 415

Ser Cys Lys Pro Gly Val Met Gly Asp Lys Cys Asp Arg Cys Gln Pro
 420 425 430

Gly Phe His Ser Leu Thr Glu Ala Gly Cys Arg Pro Cys Ser Cys Asp
 435 440 445

Pro Ser Gly Ser Ile Asp Glu Cys Asn Val Glu Thr Gly Arg Cys Val
450 455 460

Cys Lys Asp Asn Val Glu Gly Phe Asn Cys Glu Arg Cys Lys Pro Gly
465 470 475 480

Phe Phe Asn Leu Glu Ser Ser Asn Pro Arg Gly Cys Thr Pro Cys Phe
485 490 495

Cys Phe Gly His Ser Ser Val Cys Thr Asn Ala Val Gly Tyr Ser Val
500 505 510

Tyr Ser Ile Ser Ser Thr Phe Gln Ile Asp Glu Asp Gly Trp Arg Ala
515 520 525

Glu Gln Arg Asp Gly Ser Glu Ala Ser Leu Glu Trp Ser Ser Glu Arg
530 535 540

Gln Asp Ile Ala Val Ile Ser Asp Ser Tyr Phe Pro Arg Tyr Phe Ile
545 550 555 560

Ala Pro Ala Lys Phe Leu Gly Lys Gln Val Leu Ser Tyr Gly Gln Asn
565 570 575

Leu Ser Phe Ser Phe Arg Val Asp Arg Arg Asp Thr Arg Leu Ser Ala
580 585 590

Glu Asp Leu Val Leu Glu Gly Ala Gly Leu Arg Val Ser Val Pro Leu
595 600 605

Ile Ala Gln Gly Asn Ser Tyr Pro Ser Glu Thr Thr Val Lys Tyr Val
610 615 620

Phe Arg Leu His Glu Ala Thr Asp Tyr Pro Trp Arg Pro Ala Leu Thr
625 630 635 640

Pro Phe Glu Phe Gln Lys Leu Leu Asn Asn Leu Thr Ser Ile Lys Ile
645 650 655

Arg Gly Thr Tyr Ser Glu Arg Ser Ala Gly Tyr Leu Asp Asp Val Thr
660 665 670

Leu Ala Ser Ala Arg Pro Gly Pro Gly Val Pro Ala Thr Trp Val Glu

675	680	685
Ser Cys Thr Cys Pro Val Gly Tyr Gly Gly Gln Phe Cys Glu Met Cys		
690	695	700
Leu Ser Gly Tyr Arg Arg Glu Thr Pro Asn Leu Gly Pro Tyr Ser Pro		
705	710	715 720
Cys Val Leu Cys Ala Cys Asn Gly His Ser Glu Thr Cys Asp Pro Glu		
	725	730 735
Thr Gly Val Cys Asn Cys Arg Asp Asn Thr Ala Gly Pro His Cys Glu		
	740	745 750
Lys Cys Ser Asp Gly Tyr Tyr Gly Asp Ser Thr Ala Gly Thr Ser Ser		
	755	760 765
Asp Cys Gln Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Val Val		
	770	775 780
Pro Lys Thr Lys Glu Val Val Cys Thr Asn Cys Pro Thr Gly Thr Thr		
	785	790 795 800
Gly Lys Arg Cys Glu Leu Cys Asp Asp Gly Tyr Phe Gly Asp Pro Leu		
	805	810 815
Gly Arg Asn Gly Pro Val Arg Leu Cys Arg Leu Cys Gln Cys Ser Asp		
	820	825 830
Asn Ile Asp Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly Glu		
	835	840 845
Cys Leu Lys Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg Cys		
	850	855 860
Lys Asp Gly Phe Phe Gly Asn Pro Leu Ala Pro Asn Pro Ala Asp Lys		
	865	870 875 880
Cys Lys Ala Cys Asn Cys Asn Pro Tyr Gly Thr Met Lys Gln Gln Ser		
	885	890 895
Ser Cys Asn Pro Val Thr Gly Gln Cys Glu Cys Leu Pro His Val Thr		
	900	905 910

Gly Gln Asp Cys Gly Ala Cys Asp Pro Gly Phe Tyr Asn Leu Gln Ser
 915 920 925

Gly Gln Gly Cys Glu Arg Cys Asp Cys His Ala Leu Gly Ser Thr Asn
 930 935 940

Gly Gln Cys Asp Ile Arg Thr Gly Gln Cys Glu Cys Gln Pro Gly Ile
 945 950 955 960

Thr Gly Gln His Cys Glu Arg Cys Glu Val Asn His Phe Gly Phe Gly
 965 970 975

Pro Glu Gly Cys Lys Pro Cys Asp Cys His Pro Glu Gly Ser Leu Ser
 980 985 990

Leu Gln Cys Lys Asp Asp Gly Arg Cys Glu Cys Arg Glu Gly Phe Val
 995 1000 1005

Gly Asn Arg Cys Asp Gln Cys Glu Glu Asn Tyr Phe Tyr Asn Arg
 1010 1015 1020

Ser Trp Pro Gly Cys Gln Glu Cys Pro Ala Cys Tyr Arg Leu Val
 1025 1030 1035

Lys Asp Lys Val Ala Asp His Arg Val Lys Leu Gln Glu Leu Glu
 1040 1045 1050

Ser Leu Ile Ala Asn Leu Gly Thr Gly Asp Glu Met Val Thr Asp
 1055 1060 1065

Gln Ala Phe Glu Asp Arg Leu Lys Glu Ala Glu Arg Glu Val Met
 1070 1075 1080

Asp Leu Leu Arg Glu Ala Gln Asp Val Lys Asp Val Asp Gln Asn
 1085 1090 1095

Leu Met Asp Arg Leu Gln Arg Val Asn Asn Thr Leu Ser Ser Gln
 1100 1105 1110

Ile Ser Arg Leu Gln Asn Ile Arg Asn Thr Ile Glu Glu Thr Gly
 1115 1120 1125

Asn Leu Ala Glu Gln Ala Arg	Ala His Val Glu Asn Thr Glu Arg
1130	1135 1140
Leu Ile Glu Ile Ala Ser Arg	Glu Leu Glu Lys Ala Lys Val Ala
1145	1150 1155
Ala Ala Asn Val Ser Val Thr	Gln Pro Glu Ser Thr Gly Asp Pro
1160	1165 1170
Asn Asn Met Thr Leu Leu Ala	Glu Glu Ala Arg Lys Leu Ala Glu
1175	1180 1185
Arg His Lys Gln Glu Ala Asp	Asp Ile Val Arg Val Ala Lys Thr
1190	1195 1200
Ala Asn Asp Thr Ser Thr Glu	Ala Tyr Asn Leu Leu Leu Arg Thr
1205	1210 1215
Leu Ala Gly Glu Asn Gln Thr	Ala Phe Glu Ile Glu Glu Leu Asn
1220	1225 1230
Arg Lys Tyr Glu Gln Ala Lys	Asn Ile Ser Gln Asp Leu Glu Lys
1235	1240 1245
Gln Ala Ala Arg Val His Glu	Glu Ala Lys Arg Ala Gly Asp Lys
1250	1255 1260
Ala Val Glu Ile Tyr Ala Ser	Val Ala Gln Leu Ser Pro Leu Asp
1265	1270 1275
Ser Glu Thr Leu Glu Asn Glu	Ala Asn Asn Ile Lys Met Glu Ala
1280	1285 1290
Glu Asn Leu Glu Gln Leu Ile	Asp Gln Lys Leu Lys Asp Tyr Glu
1295	1300 1305
Asp Leu Arg Glu Asp Met Arg	Gly Lys Glu Leu Glu Val Lys Asn
1310	1315 1320
Leu Leu Glu Lys Gly Lys Thr	Glu Gln Gln Thr Ala Asp Gln Leu
1325	1330 1335

Leu Ala Arg Ala Asp Ala Ala Lys Ala Leu Ala Glu Glu Ala Ala
 1340 1345 1350

Lys Lys Gly Arg Asp Thr Leu Gln Glu Ala Asn Asp Ile Leu Asn
 1355 1360 1365

Asn Leu Lys Asp Phe Asp Arg Arg Val Asn Asp Asn Lys Thr Ala
 1370 1375 1380

Ala Glu Glu Ala Leu Arg Lys Ile Pro Ala Ile Asn Gln Thr Ile
 1385 1390 1395

Thr Glu Ala Asn Glu Lys Thr Arg Glu Ala Gln Gln Ala Leu Gly
 1400 1405 1410

Ser Ala Ala Ala Asp Ala Thr Glu Ala Lys Asn Lys Ala His Glu
 1415 1420 1425

Ala Glu Arg Ile Ala Ser Ala Val Gln Lys Asn Ala Thr Ser Thr
 1430 1435 1440

Lys Ala Glu Ala Glu Arg Thr Phe Ala Glu Val Thr Asp Leu Asp
 1445 1450 1455

Asn Glu Val Asn Asn Met Leu Lys Gln Leu Gln Glu Ala Glu Lys
 1460 1465 1470

Glu Leu Lys Arg Lys Gln Asp Asp Ala Asp Gln Asp Met Met Met
 1475 1480 1485

Ala Gly Met Ala Ser Gln Ala Ala Gln Glu Ala Glu Ile Asn Ala
 1490 1495 1500

Arg Lys Ala Lys Asn Ser Val Thr Ser Leu Leu Ser Ile Ile Asn
 1505 1510 1515

Asp Leu Leu Glu Gln Leu Gly Gln Leu Asp Thr Val Asp Leu Asn
 1520 1525 1530

Lys Leu Asn Glu Ile Glu Gly Thr Leu Asn Lys Ala Lys Asp Glu
 1535 1540 1545

Met Lys Val Ser Asp Leu Asp Arg Lys Val Ser Asp Leu Glu Asn

1550 1555 1560

Glu Ala Lys Lys Gln Glu Ala Ala Ile Met Asp Tyr Asn Arg Asp
1565 1570 1575

Ile Glu Glu Ile Met Lys Asp Ile Arg Asn Leu Glu Asp Ile Arg
1580 1585 1590

Lys Thr Leu Pro Ser Gly Cys Phe Asn Thr Pro Ser Ile Glu Lys
1595 1600 1605

Pro

<210> 186
<211> 1408
<212> PRT
<213> Homo sapiens

<400> 186

Met Lys Ala Pro Ala Val Leu Ala Pro Gly Ile Leu Val Leu Leu Phe
1 5 10 15

Thr Leu Val Gln Arg Ser Asn Gly Glu Cys Lys Glu Ala Leu Ala Lys
20 25 30

Ser Glu Met Asn Val Asn Met Lys Tyr Gln Leu Pro Asn Phe Thr Ala
35 40 45

Glu Thr Pro Ile Gln Asn Val Ile Leu His Glu His His Ile Phe Leu
50 55 60

Gly Ala Thr Asn Tyr Ile Tyr Val Leu Asn Glu Glu Asp Leu Gln Lys
65 70 75 80

Val Ala Glu Tyr Lys Thr Gly Pro Val Leu Glu His Pro Asp Cys Phe
85 90 95

Pro Cys Gln Asp Cys Ser Ser Lys Ala Asn Leu Ser Gly Gly Val Trp
100 105 110

Lys Asp Asn Ile Asn Met Ala Leu Val Val Asp Thr Tyr Tyr Asp Asp
115 120 125

Gln Leu Ile Ser Cys Gly Ser Val Asn Arg Gly Thr Cys Gln Arg His
 130 135 140

Val Phe Pro His Asn His Thr Ala Asp Ile Gln Ser Glu Val His Cys
 145 150 155 160

Ile Phe Ser Pro Gln Ile Glu Glu Pro Ser Gln Cys Pro Asp Cys Val
 165 170 175

Val Ser Ala Leu Gly Ala Lys Val Leu Ser Ser Val Lys Asp Arg Phe
 180 185 190

Ile Asn Phe Phe Val Gly Asn Thr Ile Asn Ser Ser Tyr Phe Pro Asp
 195 200 205

His Pro Leu His Ser Ile Ser Val Arg Arg Leu Lys Glu Thr Lys Asp
 210 215 220

Gly Phe Met Phe Leu Thr Asp Gln Ser Tyr Ile Asp Val Leu Pro Glu
 225 230 235 240

Phe Arg Asp Ser Tyr Pro Ile Lys Tyr Val His Ala Phe Glu Ser Asn
 245 250 255

Asn Phe Ile Tyr Phe Leu Thr Val Gln Arg Glu Thr Leu Asp Ala Gln
 260 265 270

Thr Phe His Thr Arg Ile Ile Arg Phe Cys Ser Ile Asn Ser Gly Leu
 275 280 285

His Ser Tyr Met Glu Met Pro Leu Glu Cys Ile Leu Thr Glu Lys Arg
 290 295 300

Lys Lys Arg Ser Thr Lys Lys Glu Val Phe Asn Ile Leu Gln Ala Ala
 305 310 315 320

Tyr Val Ser Lys Pro Gly Ala Gln Leu Ala Arg Gln Ile Gly Ala Ser
 325 330 335

Leu Asn Asp Asp Ile Leu Phe Gly Val Phe Ala Gln Ser Lys Pro Asp
 340 345 350

Ser Ala Glu Pro Met Asp Arg Ser Ala Met Cys Ala Phe Pro Ile Lys
 355 360 365

Tyr Val Asn Asp Phe Phe Asn Lys Ile Val Asn Lys Asn Asn Val Arg
 370 375 380

Cys Leu Gln His Phe Tyr Gly Pro Asn His Glu His Cys Phe Asn Arg
 385 390 395 400

Thr Leu Leu Arg Asn Ser Ser Gly Cys Glu Ala Arg Arg Asp Glu Tyr
 405 410 415

Arg Thr Glu Phe Thr Thr Ala Leu Gln Arg Val Asp Leu Phe Met Gly
 420 425 430

Gln Phe Ser Glu Val Leu Leu Thr Ser Ile Ser Thr Phe Ile Lys Gly
 435 440 445

Asp Leu Thr Ile Ala Asn Leu Gly Thr Ser Glu Gly Arg Phe Met Gln
 450 455 460

Val Val Val Ser Arg Ser Gly Pro Ser Thr Pro His Val Asn Phe Leu
 465 470 475 480

Leu Asp Ser His Pro Val Ser Pro Glu Val Ile Val Glu His Thr Leu
 485 490 495

Asn Gln Asn Gly Tyr Thr Leu Val Ile Thr Gly Lys Lys Ile Thr Lys
 500 505 510

Ile Pro Leu Asn Gly Leu Gly Cys Arg His Phe Gln Ser Cys Ser Gln
 515 520 525

Cys Leu Ser Ala Pro Pro Phe Val Gln Cys Gly Trp Cys His Asp Lys
 530 535 540

Cys Val Arg Ser Glu Glu Cys Leu Ser Gly Thr Trp Thr Gln Gln Ile
 545 550 555 560

Cys Leu Pro Ala Ile Tyr Lys Val Phe Pro Asn Ser Ala Pro Leu Glu
 565 570 575

Gly Gly Thr Arg Leu Thr Ile Cys Gly Trp Asp Phe Gly Phe Arg Arg

580	585	590
Asn Asn Lys Phe Asp Leu Lys Lys Thr Arg Val Leu Leu Gly Asn Glu		
595	600	605
Ser Cys Thr Leu Thr Leu Ser Glu Ser Thr Met Asn Thr Leu Lys Cys		
610	615	620
Thr Val Gly Pro Ala Met Asn Lys His Phe Asn Met Ser Ile Ile Ile		
625	630	635 640
Ser Asn Gly His Gly Thr Thr Gln Tyr Ser Thr Phe Ser Tyr Val Asp		
645	650	655
Pro Val Ile Thr Ser Ile Ser Pro Lys Tyr Gly Pro Met Ala Gly Gly		
660	665	670
Thr Leu Leu Thr Leu Thr Gly Asn Tyr Leu Asn Ser Gly Asn Ser Arg		
675	680	685
His Ile Ser Ile Gly Gly Lys Thr Cys Thr Leu Lys Ser Val Ser Asn		
690	695	700
Ser Ile Leu Glu Cys Tyr Thr Pro Ala Gln Thr Ile Ser Thr Glu Phe		
705	710	715 720
Ala Val Lys Leu Lys Ile Asp Leu Ala Asn Arg Glu Thr Ser Ile Phe		
725	730	735
Ser Tyr Arg Glu Asp Pro Ile Val Tyr Glu Ile His Pro Thr Lys Ser		
740	745	750
Phe Ile Ser Thr Trp Trp Lys Glu Pro Leu Asn Ile Val Ser Phe Leu		
755	760	765
Phe Cys Phe Ala Ser Gly Gly Ser Thr Ile Thr Gly Val Gly Lys Asn		
770	775	780
Leu Asn Ser Val Ser Val Pro Arg Met Val Ile Asn Val His Glu Ala		
785	790	795 800
Gly Arg Asn Phe Thr Val Ala Cys Gln His Arg Ser Asn Ser Glu Ile		
805	810	815

Ile Cys Cys Thr Thr Pro Ser Leu Gln Gln Leu Asn Leu Gln Leu Pro
820 825 830

Leu Lys Thr Lys Ala Phe Phe Met Leu Asp Gly Ile Leu Ser Lys Tyr
835 840 845

Phe Asp Leu Ile Tyr Val His Asn Pro Val Phe Lys Pro Phe Glu Lys
850 855 860

Pro Val Met Ile Ser Met Gly Asn Glu Asn Val Leu Glu Ile Lys Gly
865 870 875 880

Asn Asp Ile Asp Pro Glu Ala Val Lys Gly Glu Val Leu Lys Val Gly
885 890 895

Asn Lys Ser Cys Glu Asn Ile His Leu His Ser Glu Ala Val Leu Cys
900 905 910

Thr Val Pro Asn Asp Leu Leu Lys Leu Asn Ser Glu Leu Asn Ile Glu
915 920 925

Trp Lys Gln Ala Ile Ser Ser Thr Val Leu Gly Lys Val Ile Val Gln
930 935 940

Pro Asp Gln Asn Phe Thr Gly Leu Ile Ala Gly Val Val Ser Ile Ser
945 950 955 960

Thr Ala Leu Leu Leu Leu Leu Gly Phe Phe Leu Trp Leu Lys Lys Arg
965 970 975

Lys Gln Ile Lys Asp Leu Gly Ser Glu Leu Val Arg Tyr Asp Ala Arg
980 985 990

Val His Thr Pro His Leu Asp Arg Leu Val Ser Ala Arg Ser Val Ser
995 1000 1005

Pro Thr Thr Glu Met Val Ser Asn Glu Ser Val Asp Tyr Arg Ala
1010 1015 1020

Thr Phe Pro Glu Asp Gln Phe Pro Asn Ser Ser Gln Asn Gly Ser
1025 1030 1035

297

Asp Lys Glu Tyr Tyr Ser Val His Asn Lys Thr Gly Ala Lys Leu
1250 1255 1260

Pro Val Lys Trp Met Ala Leu Glu Ser Leu Gln Thr Gln Lys Phe
1265 1270 1275

Thr Thr Lys Ser Asp Val Trp Ser Phe Gly Val Val Leu Trp Glu
1280 1285 1290

Leu Met Thr Arg Gly Ala Pro Pro Tyr Pro Asp Val Asn Thr Phe
1295 1300 1305

Asp Ile Thr Val Tyr Leu Leu Gln Gly Arg Arg Leu Leu Gln Pro
1310 1315 1320

Glu Tyr Cys Pro Asp Pro Leu Tyr Glu Val Met Leu Lys Cys Trp
1325 1330 1335

His Pro Lys Ala Glu Met Arg Pro Ser Phe Ser Glu Leu Val Ser
1340 1345 1350

Arg Ile Ser Ala Ile Phe Ser Thr Phe Ile Gly Glu His Tyr Val
1355 1360 1365

His Val Asn Ala Thr Tyr Val Asn Val Lys Cys Val Ala Pro Tyr
1370 1375 1380

Pro Ser Leu Leu Ser Ser Glu Asp Asn Ala Asp Asp Glu Val Asp
1385 1390 1395

Thr Arg Pro Ala Ser Phe Trp Glu Thr Ser
1400 1405

<210> 187

<211> 577

<212> PRT

<213> Homo sapiens

<400> 187

Met Pro Lys Thr Ile Ser Val Arg Val Thr Thr Met Asp Ala Glu Leu
1 5 10 15

Glu Phe Ala Ile Gln Pro Asn Thr Thr Gly Lys Gln Leu Phe Asp Gln
20 25 30

Val Val Lys Thr Ile Gly Leu Arg Glu Val Trp Phe Phe Gly Leu Gln
 35 40 45

Tyr Gln Asp Thr Lys Gly Phe Ser Thr Trp Leu Lys Leu Asn Lys Lys
 50 55 60

Val Thr Ala Gln Asp Val Arg Lys Glu Ser Pro Leu Leu Phe Lys Phe
 65 70 75 80

Arg Ala Lys Phe Tyr Pro Glu Asp Val Ser Glu Glu Leu Ile Gln Asp
 85 90 95

Ile Thr Gln Arg Leu Phe Phe Leu Gln Val Lys Glu Gly Ile Leu Asn
 100 105 110

Asp Asp Ile Tyr Cys Pro Pro Glu Thr Ala Val Leu Leu Ala Ser Tyr
 115 120 125

Ala Val Gln Ser Lys Tyr Gly Asp Phe Asn Lys Glu Val His Lys Ser
 130 135 140

Gly Tyr Leu Ala Gly Asp Lys Leu Leu Pro Gln Arg Val Leu Glu Gln
 145 150 155 160

His Lys Leu Asn Lys Asp Gln Trp Glu Glu Arg Ile Gln Val Trp His
 165 170 175

Glu Glu His Arg Gly Met Leu Arg Glu Asp Ala Val Leu Glu Tyr Leu
 180 185 190

Lys Ile Ala Gln Asp Leu Glu Met Tyr Gly Val Asn Tyr Phe Ser Ile
 195 200 205

Lys Asn Lys Lys Gly Ser Glu Leu Trp Leu Gly Val Asp Ala Leu Gly
 210 215 220

Leu Asn Ile Tyr Glu Gln Asn Asp Arg Leu Thr Pro Lys Ile Gly Phe
 225 230 235 240

Pro Trp Ser Glu Ile Arg Asn Ile Ser Phe Asn Asp Lys Lys Phe Val
 245 250 255

Ile Lys Pro Ile Asp Lys Lys Ala Pro Asp Phe Val Phe Tyr Ala Pro
 260 265 270

Arg Leu Arg Ile Asn Lys Arg Ile Leu Ala Leu Cys Met Gly Asn His
 275 280 285

Glu Leu Tyr Met Arg Arg Arg Lys Pro Asp Thr Ile Glu Val Gln Gln
 290 295 300

Met Lys Ala Gln Ala Arg Glu Glu Lys His Gln Lys Gln Met Glu Arg
 305 310 315 320

Ala Met Leu Glu Asn Glu Lys Lys Lys Arg Glu Met Ala Glu Lys Glu
 325 330 335

Lys Glu Lys Ile Glu Arg Glu Lys Glu Glu Leu Met Glu Arg Leu Lys
 340 345 350

Gln Ile Glu Glu Gln Thr Lys Lys Ala Gln Gln Glu Leu Glu Glu Gln
 355 360 365

Thr Arg Arg Ala Leu Glu Leu Glu Gln Glu Arg Lys Arg Ala Gln Ser
 370 375 380

Glu Ala Glu Lys Leu Ala Lys Glu Arg Gln Glu Ala Glu Glu Ala Lys
 385 390 395 400

Glu Ala Leu Leu Gln Ala Ser Arg Asp Gln Lys Lys Thr Gln Glu Gln
 405 410 415

Leu Ala Leu Glu Met Ala Glu Leu Thr Ala Arg Ile Ser Gln Leu Glu
 420 425 430

Met Ala Arg Gln Lys Lys Glu Ser Glu Ala Val Glu Trp Gln Gln Lys
 435 440 445

Ala Gln Met Val Gln Glu Asp Leu Glu Lys Thr Arg Ala Glu Leu Lys
 450 455 460

Thr Ala Met Ser Thr Pro His Val Ala Glu Pro Ala Glu Asn Glu Gln
 465 470 475 480

Asp Glu Gln Asp Glu Asn Gly Ala Glu Ala Ser Ala Asp Leu Arg Ala
 485 490 495

Asp Ala Met Ala Lys Asp Arg Ser Glu Glu Glu Arg Thr Thr Glu Ala
 500 505 510

Glu Lys Asn Glu Arg Val Gln Lys His Leu Lys Ala Leu Thr Ser Glu
 515 520 525

Leu Ala Asn Ala Arg Asp Glu Ser Lys Lys Thr Ala Asn Asp Met Ile
 530 535 540

His Ala Glu Asn Met Arg Leu Gly Arg Asp Lys Tyr Lys Thr Leu Arg
 545 550 555 560

Gln Ile Arg Gln Gly Asn Thr Lys Gln Arg Ile Asp Glu Phe Glu Ser
 565 570 575

Met

<210> 188
 <211> 2058
 <212> PRT
 <213> Homo sapiens

<400> 188

Met Asp Asn Phe Phe Thr Glu Gly Thr Arg Val Trp Leu Arg Glu Asn
 1 5 10 15

Gly Gln His Phe Pro Ser Thr Val Asn Ser Cys Ala Glu Gly Ile Val
 20 25 30

Val Phe Arg Thr Asp Tyr Gly Gln Val Phe Thr Tyr Lys Gln Ser Thr
 35 40 45

Ile Thr His Gln Lys Val Thr Ala Met His Pro Thr Asn Glu Glu Gly
 50 55 60

Val Asp Asp Met Ala Ser Leu Thr Glu Leu His Gly Gly Ser Ile Met
 65 70 75 80

Tyr Asn Leu Phe Gln Arg Tyr Lys Arg Asn Gln Ile Tyr Thr Tyr Ile
 85 90 95

Gly Ser Ile Leu Ala Ser Val Asn Pro Tyr Gln Pro Ile Ala Gly Leu
 100 105 110

Tyr Glu Pro Ala Thr Met Glu Gln Tyr Ser Arg Arg His Leu Gly Glu
 115 120 125

Leu Pro Pro His Ile Phe Ala Ile Ala Asn Glu Cys Tyr Arg Cys Leu
 130 135 140

Trp Lys Arg Tyr Asp Asn Gln Cys Ile Leu Ile Ser Gly Glu Ser Gly
 145 150 155 160

Ala Gly Lys Thr Glu Ser Thr Lys Leu Ile Leu Lys Phe Leu Ser Val
 165 170 175

Ile Ser Gln Gln Ser Leu Glu Leu Ser Leu Lys Glu Lys Thr Ser Cys
 180 185 190

Val Glu Arg Ala Ile Leu Glu Ser Ser Pro Ile Met Glu Ala Phe Gly
 195 200 205

Asn Ala Lys Thr Val Tyr Asn Asn Asn Ser Ser Arg Phe Gly Lys Phe
 210 215 220

Val Gln Leu Asn Ile Cys Gln Lys Gly Asn Ile Gln Gly Gly Arg Ile
 225 230 235 240

Val Asp Tyr Leu Leu Glu Lys Asn Arg Val Val Arg Gln Asn Pro Gly
 245 250 255

Glu Arg Asn Tyr His Ile Phe Tyr Ala Leu Leu Ala Gly Leu Glu His
 260 265 270

Glu Glu Arg Glu Glu Phe Tyr Leu Ser Thr Pro Glu Asn Tyr His Tyr
 275 280 285

Leu Asn Gln Ser Gly Cys Val Glu Asp Lys Thr Ile Ser Asp Gln Glu
 290 295 300

Ser Phe Arg Glu Val Ile Thr Ala Met Asp Val Met Gln Phe Ser Lys
 305 310 315 320

Glu Glu Val Arg Glu Val Ser Arg Leu Leu Ala Gly Ile Leu His Leu
325 330 335

Gly Asn Ile Glu Phe Ile Thr Ala Gly Gly Ala Gln Val Ser Phe Lys
340 345 350

Thr Ala Leu Gly Arg Ser Ala Glu Leu Leu Gly Leu Asp Pro Thr Gln
355 360 365

Leu Thr Asp Ala Leu Thr Gln Arg Ser Met Phe Leu Arg Gly Glu Glu
370 375 380

Ile Leu Thr Pro Leu Asn Val Gln Gln Ala Val Asp Ser Arg Asp Ser
385 390 395 400

Leu Ala Met Ala Leu Tyr Ala Cys Cys Phe Glu Trp Val Ile Lys Lys
405 410 415

Ile Asn Ser Arg Ile Lys Gly Asn Glu Asp Phe Lys Ser Ile Gly Ile
420 425 430

Leu Asp Ile Phe Gly Phe Glu Asn Phe Glu Val Asn His Phe Glu Gln
435 440 445

Phe Asn Ile Asn Tyr Ala Asn Glu Lys Leu Gln Glu Tyr Phe Asn Lys
450 455 460

His Ile Phe Ser Leu Glu Gln Leu Glu Tyr Ser Arg Glu Gly Leu Val
465 470 475 480

Trp Glu Asp Ile Asp Trp Ile Asp Asn Gly Glu Cys Leu Asp Leu Ile
485 490 495

Glu Lys Lys Leu Gly Leu Leu Ala Leu Ile Asn Glu Glu Ser His Phe
500 505 510

Pro Gln Ala Thr Asp Ser Thr Leu Leu Glu Lys Leu His Ser Gln His
515 520 525

Ala Asn Asn His Phe Tyr Val Lys Pro Arg Val Ala Val Asn Asn Phe
530 535 540

Gly Val Lys His Tyr Ala Gly Glu Val Gln Tyr Asp Val Arg Gly Ile
545 550 555 560

Leu Glu Lys Asn Arg Asp Thr Phe Arg Asp Asp Leu Leu Asn Leu Leu
565 570 575

Arg Glu Ser Arg Phe Asp Phe Ile Tyr Asp Leu Phe Glu His Val Ser
580 585 590

Ser Arg Asn Asn Gln Asp Thr Leu Lys Cys Gly Ser Lys His Arg Arg
595 600 605

Pro Thr Val Ser Ser Gln Phe Lys Asp Ser Leu His Ser Leu Met Ala
610 615 620

Thr Leu Ser Ser Ser Asn Pro Phe Phe Val Arg Cys Ile Lys Pro Asn
625 630 635 640

Met Gln Lys Met Pro Asp Gln Phe Asp Gln Ala Val Val Leu Asn Gln
645 650 655

Leu Arg Tyr Ser Gly Met Leu Glu Thr Val Arg Ile Arg Lys Ala Gly
660 665 670

Tyr Ala Val Arg Arg Pro Phe Gln Asp Phe Tyr Lys Arg Tyr Lys Val
675 680 685

Leu Met Arg Asn Leu Ala Leu Pro Glu Asp Val Arg Gly Lys Cys Thr
690 695 700

Ser Leu Leu Gln Leu Tyr Asp Ala Ser Asn Ser Glu Trp Gln Leu Gly
705 710 715 720

Lys Thr Lys Val Phe Leu Arg Glu Ser Leu Glu Gln Lys Leu Glu Lys
725 730 735

Arg Arg Glu Glu Glu Val Ser His Ala Ala Met Val Ile Arg Ala His
740 745 750

Val Leu Gly Phe Leu Ala Arg Lys Gln Tyr Arg Lys Val Leu Tyr Cys
755 760 765

Val Val Ile Ile Gln Lys Asn Tyr Arg Ala Phe Leu Leu Arg Arg Arg

770	775	780
Phe Leu His Leu Lys Lys Ala Ala Ile Val Phe Gln Lys Gln Leu Arg		
785	790	795 800
Gly Gln Ile Ala Arg Arg Val Tyr Arg Gln Leu Leu Ala Glu Lys Arg		
	805	810 815
Glu Gln Glu Glu Lys Lys Lys Gln Glu Glu Glu Lys Lys Lys Arg		
	820	825 830
Glu Glu Glu Glu Arg Glu Arg Glu Arg Glu Arg Arg Glu Ala Glu Leu		
	835	840 845
Arg Ala Gln Gln Glu Glu Glu Thr Arg Lys Gln Gln Glu Leu Glu Ala		
	850	855 860
Leu Gln Lys Ser Gln Lys Glu Ala Glu Leu Thr Arg Glu Leu Glu Lys		
865	870	875 880
Gln Lys Glu Asn Lys Gln Val Glu Glu Ile Leu Arg Leu Glu Lys Glu		
	885	890 895
Ile Glu Asp Leu Gln Arg Met Lys Glu Gln Gln Glu Leu Ser Leu Thr		
	900	905 910
Glu Ala Ser Leu Gln Lys Leu Gln Glu Arg Arg Asp Gln Glu Leu Arg		
	915	920 925
Arg Leu Glu Glu Glu Ala Cys Arg Ala Ala Gln Glu Phe Leu Glu Ser		
	930	935 940
Leu Asn Phe Asp Glu Ile Asp Glu Cys Val Arg Asn Ile Glu Arg Ser		
945	950	955 960
Leu Ser Val Gly Ser Glu Phe Ser Ser Glu Leu Ala Glu Ser Ala Cys		
	965	970 975
Glu Glu Lys Pro Asn Phe Asn Phe Ser Gln Pro Tyr Pro Glu Glu Glu		
	980	985 990
Val Asp Glu Gly Phe Glu Ala Asp Asp Asp Ala Phe Lys Asp Ser Pro		
	995	1000 1005

Asn Pro	Ser Glu His Gly His	Ser Asp Gln Arg Thr	Ser Gly Ile
1010	1015	1020	
Arg Thr	Ser Asp Asp Ser Ser	Glu Glu Asp Pro Tyr	Met Asn Asp
1025	1030	1035	
Thr Val	Val Pro Thr Ser Pro	Ser Ala Asp Ser Thr	Val Leu Leu
1040	1045	1050	
Ala Pro	Ser Val Gln Asp Ser	Gly Ser Leu His Asn	Ser Ser Ser
1055	1060	1065	
Gly Glu	Ser Thr Tyr Cys Met	Pro Gln Asn Ala Gly	Asp Leu Pro
1070	1075	1080	
Ser Pro	Asp Gly Asp Tyr Asp	Tyr Asp Gln Asp Asp	Tyr Glu Asp
1085	1090	1095	
Gly Ala	Ile Thr Ser Gly Ser	Ser Val Thr Phe Ser	Asn Ser Tyr
1100	1105	1110	
Gly Ser	Gln Trp Ser Pro Asp	Tyr Arg Cys Ser Val	Gly Thr Tyr
1115	1120	1125	
Asn Ser	Ser Gly Ala Tyr Arg	Phe Ser Ser Glu Gly	Ala Gln Ser
1130	1135	1140	
Ser Phe	Glu Asp Ser Glu Glu	Asp Phe Asp Ser Arg	Phe Asp Thr
1145	1150	1155	
Asp Asp	Glu Leu Ser Tyr Arg	Arg Asp Ser Val Tyr	Ser Cys Val
1160	1165	1170	
Thr Leu	Pro Tyr Phe His Ser	Phe Leu Tyr Met Lys	Gly Gly Leu
1175	1180	1185	
Met Asn	Ser Trp Lys Arg Arg	Trp Cys Val Leu Lys	Asp Glu Thr
1190	1195	1200	
Phe Leu	Trp Phe Arg Ser Lys	Gln Glu Ala Leu Lys	Gln Gly Trp
1205	1210	1215	

Leu His	Lys Lys Gly Gly Gly	Ser Ser Thr Leu Ser	Arg Arg Asn
1220	1225	1230	
Trp Lys	Lys Arg Trp Phe Val	Leu Arg Gln Ser Lys	Leu Met Tyr
1235	1240	1245	
Phe Glu	Asn Asp Ser Glu Glu	Lys Leu Lys Gly Thr	Val Glu Val
1250	1255	1260	
Arg Thr	Ala Lys Glu Ile Ile	Asp Asn Thr Thr Lys	Glu Asn Gly
1265	1270	1275	
Ile Asp	Ile Ile Met Ala Asp	Arg Thr Phe His Leu	Ile Ala Glu
1280	1285	1290	
Ser Pro	Glu Asp Ala Ser Gln	Trp Phe Ser Val Leu	Ser Gln Val
1295	1300	1305	
His Ala	Ser Thr Asp Gln Glu	Ile Gln Glu Met His	Asp Glu Gln
1310	1315	1320	
Ala Asn	Pro Gln Asn Ala Val	Gly Thr Leu Asp Val	Gly Leu Ile
1325	1330	1335	
Asp Ser	Val Cys Ala Ser Asp	Ser Pro Asp Arg Pro	Asn Ser Phe
1340	1345	1350	
Val Ile	Ile Thr Ala Asn Arg	Val Leu His Cys Asn	Ala Asp Thr
1355	1360	1365	
Pro Glu	Glu Met His His Trp	Ile Thr Leu Leu Gln	Arg Ser Lys
1370	1375	1380	
Gly Asp	Thr Arg Val Glu Gly	Gln Glu Phe Ile Val	Arg Gly Trp
1385	1390	1395	
Leu His	Lys Glu Val Lys Asn	Ser Pro Lys Met Ser	Ser Leu Lys
1400	1405	1410	
Leu Lys	Lys Arg Trp Phe Val	Leu Thr His Asn Ser	Leu Asp Tyr
1415	1420	1425	

Tyr	Lys	Ser	Ser	Glu	Lys	Asn	Ala	Leu	Lys	Leu	Gly	Thr	Leu	Val
1430						1435					1440			
Leu	Asn	Ser	Leu	Cys	Ser	Val	Val	Pro	Pro	Asp	Glu	Lys	Ile	Phe
1445						1450					1455			
Lys	Glu	Thr	Gly	Tyr	Trp	Asn	Val	Thr	Val	Tyr	Gly	Arg	Lys	His
1460						1465					1470			
Cys	Tyr	Arg	Leu	Tyr	Thr	Lys	Leu	Leu	Asn	Glu	Ala	Thr	Arg	Trp
1475						1480					1485			
Ser	Ser	Ala	Ile	Gln	Asn	Val	Thr	Asp	Thr	Lys	Ala	Pro	Ile	Asp
1490						1495					1500			
Thr	Pro	Thr	Gln	Gln	Leu	Ile	Gln	Asp	Ile	Lys	Glu	Asn	Cys	Leu
1505						1510					1515			
Asn	Ser	Asp	Val	Val	Glu	Gln	Ile	Tyr	Lys	Arg	Asn	Pro	Ile	Leu
1520						1525					1530			
Arg	Tyr	Thr	His	His	Pro	Leu	His	Ser	Pro	Leu	Leu	Pro	Leu	Pro
1535						1540					1545			
Tyr	Gly	Asp	Ile	Asn	Leu	Asn	Leu	Leu	Lys	Asp	Lys	Gly	Tyr	Thr
1550						1555					1560			
Thr	Leu	Gln	Asp	Glu	Ala	Ile	Lys	Ile	Phe	Asn	Ser	Leu	Gln	Gln
1565						1570					1575			
Leu	Glu	Ser	Met	Ser	Asp	Pro	Ile	Pro	Ile	Ile	Gln	Gly	Ile	Leu
1580						1585					1590			
Gln	Thr	Gly	His	Asp	Leu	Arg	Pro	Leu	Arg	Asp	Glu	Leu	Tyr	Cys
1595						1600					1605			
Gln	Leu	Ile	Lys	Gln	Thr	Asn	Lys	Val	Pro	His	Pro	Gly	Ser	Val
1610						1615					1620			
Gly	Asn	Leu	Tyr	Ser	Trp	Gln	Ile	Leu	Thr	Cys	Leu	Ser	Cys	Thr
1625						1630					1635			
Phe	Leu	Pro	Ser	Arg	Gly	Ile	Leu	Lys	Tyr	Leu	Lys	Phe	His	Leu

1640	1645	1650
Lys Arg Ile Arg Glu Gln Phe Pro Gly Thr Glu Met Glu Lys Tyr 1655 1660 1665		
Ala Leu Phe Thr Tyr Glu Ser Leu Lys Lys Thr Lys Cys Arg Glu 1670 1675 1680		
Phe Val Pro Ser Arg Asp Glu Ile Glu Ala Leu Ile His Arg Gln 1685 1690 1695		
Glu Met Thr Ser Thr Val Tyr Cys His Gly Gly Gly Ser Cys Lys 1700 1705 1710		
Ile Thr Ile Asn Ser His Thr Thr Ala Gly Glu Val Val Glu Lys 1715 1720 1725		
Leu Ile Arg Gly Leu Ala Met Glu Asp Ser Arg Asn Met Phe Ala 1730 1735 1740		
Leu Phe Glu Tyr Asn Gly His Val Asp Lys Ala Ile Glu Ser Arg 1745 1750 1755		
Thr Val Val Ala Asp Val Leu Ala Lys Phe Glu Lys Leu Ala Ala 1760 1765 1770		
Thr Ser Glu Val Gly Asp Leu Pro Trp Lys Phe Tyr Phe Lys Leu 1775 1780 1785		
Tyr Cys Phe Leu Asp Thr Asp Asn Val Pro Lys Asp Ser Val Glu 1790 1795 1800		
Phe Ala Phe Met Phe Glu Gln Ala His Glu Ala Val Ile His Gly 1805 1810 1815		
His His Pro Ala Pro Glu Glu Asn Leu Gln Val Leu Ala Ala Leu 1820 1825 1830		
Arg Leu Gln Tyr Leu Gln Gly Asp Tyr Thr Leu His Ala Ala Ile 1835 1840 1845		
Pro Pro Leu Glu Glu Val Tyr Ser Leu Gln Arg Leu Lys Ala Arg 1850 1855 1860		

Ile Ser Gln Ser Thr Lys Thr Phe Thr Pro Cys Glu Arg Leu Glu
1865 1870 1875

Lys Arg Arg Thr Ser Phe Leu Glu Gly Thr Leu Arg Arg Ser Phe
1880 1885 1890

Arg Thr Gly Ser Val Val Arg Gln Lys Val Glu Glu Glu Gln Met
1895 1900 1905

Leu Asp Met Trp Ile Lys Glu Glu Val Ser Ser Ala Arg Ala Ser
1910 1915 1920

Ile Ile Asp Lys Trp Arg Lys Phe Gln Gly Met Asn Gln Glu Gln
1925 1930 1935

Ala Met Ala Lys Tyr Met Ala Leu Ile Lys Glu Trp Pro Gly Tyr
1940 1945 1950

Gly Ser Thr Leu Phe Asp Val Glu Cys Lys Glu Gly Gly Phe Pro
1955 1960 1965

Gln Glu Leu Trp Leu Gly Val Ser Ala Asp Ala Val Ser Val Tyr
1970 1975 1980

Lys Arg Gly Glu Gly Arg Pro Leu Glu Val Phe Gln Tyr Glu His
1985 1990 1995

Ile Leu Ser Phe Gly Ala Pro Leu Ala Asn Thr Tyr Lys Ile Val
2000 2005 2010

Val Asp Glu Arg Glu Leu Leu Phe Glu Thr Ser Glu Val Val Asp
2015 2020 2025

Val Ala Lys Leu Met Lys Ala Tyr Ile Ser Met Ile Val Lys Lys
2030 2035 2040

Arg Tyr Ser Thr Thr Arg Ser Ala Ser Ser Gln Gly Ser Ser Arg
2045 2050 2055

<210> 189
<211> 562
<212> PRT

<213> Homo sapiens

<400> 189

Met Val Lys Ile Val Thr Val Lys Thr Gln Ala Tyr Gln Asp Gln Lys
 1 5 10 15

Pro Gly Thr Ser Gly Leu Arg Lys Arg Val Lys Val Phe Gln Ser Ser
 20 25 30

Ala Asn Tyr Ala Glu Asn Phe Ile Gln Ser Ile Ile Ser Thr Val Glu
 35 40 45

Pro Ala Gln Arg Gln Glu Ala Thr Leu Val Val Gly Gly Asp Gly Arg
 50 55 60

Phe Tyr Met Lys Glu Ala Ile Gln Leu Ile Ala Arg Ile Ala Ala Ala
 65 70 75 80

Asn Gly Ile Gly Arg Leu Val Ile Gly Gln Asn Gly Ile Leu Ser Thr
 85 90 95

Pro Ala Val Ser Cys Ile Ile Arg Lys Ile Lys Ala Ile Gly Gly Ile
 100 105 110

Ile Leu Thr Ala Ser His Asn Pro Gly Gly Pro Asn Gly Asp Phe Gly
 115 120 125

Ile Lys Phe Asn Ile Ser Asn Gly Gly Pro Ala Pro Glu Ala Ile Thr
 130 135 140

Asp Lys Ile Phe Gln Ile Ser Lys Thr Ile Glu Glu Tyr Ala Val Cys
 145 150 155 160

Pro Asp Leu Lys Val Asp Leu Gly Val Leu Gly Lys Gln Gln Phe Asp
 165 170 175

Leu Glu Asn Lys Phe Lys Pro Phe Thr Val Glu Ile Val Asp Ser Val
 180 185 190

Glu Ala Tyr Ala Thr Met Leu Arg Ser Ile Phe Asp Phe Ser Ala Leu
 195 200 205

Lys Glu Leu Leu Ser Gly Pro Asn Arg Leu Lys Ile Cys Ile Asp Ala

210	215	220
Met His Gly Val Val Gly Pro Tyr Val Lys Lys Ile Leu Cys Glu Glu 225 230 235 240		
Leu Gly Ala Pro Ala Asn Ser Ala Val Asn Cys Val Pro Leu Glu Asp 245 250 255		
Phe Gly Gly His His Pro Asp Pro Asn Leu Thr Tyr Ala Ala Asp Leu 260 265 270		
Val Glu Thr Met Lys Ser Gly Glu His Asp Phe Gly Ala Ala Phe Asp 275 280 285		
Gly Asp Gly Asp Arg Asn Met Ile Leu Gly Lys His Gly Phe Phe Val 290 295 300		
Asn Pro Ser Asp Ser Val Ala Val Ile Ala Ala Asn Ile Phe Ser Ile 305 310 315 320		
Pro Tyr Phe Gln Gln Thr Gly Val Arg Gly Phe Ala Arg Ser Met Pro 325 330 335		
Thr Ser Gly Ala Leu Asp Arg Val Ala Ser Ala Thr Lys Ile Ala Leu 340 345 350		
Tyr Glu Thr Pro Thr Gly Trp Lys Phe Phe Gly Asn Leu Met Asp Ala 355 360 365		
Ser Lys Leu Ser Leu Cys Gly Glu Glu Ser Phe Gly Thr Gly Ser Asp 370 375 380		
His Ile Arg Glu Lys Asp Gly Leu Trp Ala Val Leu Ala Trp Leu Ser 385 390 395 400		
Ile Leu Ala Thr Arg Lys Gln Ser Val Glu Asp Ile Leu Lys Asp His 405 410 415		
Trp Gln Lys His Gly Arg Asn Phe Phe Thr Arg Tyr Asp Tyr Glu Glu 420 425 430		
Val Glu Ala Glu Gly Ala Asn Lys Met Met Lys Asp Leu Glu Ala Leu 435 440 445		

Met Phe Asp Arg Ser Phe Val Gly Lys Gln Phe Ser Ala Asn Asp Lys
 450 455 460

Val Tyr Thr Val Glu Lys Ala Asp Asn Phe Glu Tyr Ser Asp Pro Val
 465 470 475 480

Asp Gly Ser Ile Ser Arg Asn Gln Gly Leu Arg Leu Ile Phe Thr Asp
 485 490 495

Gly Ser Arg Ile Val Phe Arg Leu Ser Gly Thr Gly Ser Ala Gly Ala
 500 505 510

Thr Ile Arg Leu Tyr Ile Asp Ser Tyr Glu Lys Asp Val Ala Lys Ile
 515 520 525

Asn Gln Asp Pro Gln Val Met Leu Ala Pro Leu Ile Ser Ile Ala Leu
 530 535 540

Lys Val Ser Gln Leu Gln Glu Arg Thr Gly Arg Thr Ala Pro Thr Val
 545 550 555 560

Ile Thr

<210> 190
 <211> 204
 <212> PRT
 <213> Homo sapiens

<400> 190

Gly Glu Gly Glu Arg Pro Glu Glu Asp Ala Ala Ala Leu Glu Leu Ser
 1 5 10 15

Ser Asp Glu Ala Val Glu Val Glu Glu Val Ile Glu Glu Ser Arg Ala
 20 25 30

Glu Arg Ile Lys Arg Ser Gly Leu Arg Arg Val Asp Asp Phe Lys Lys
 35 40 45

Ala Phe Ser Lys Glu Lys Met Glu Lys Thr Lys Val Arg Thr Arg Glu
 50 55 60

Asn Leu Glu Lys Thr Arg Leu Lys Thr Lys Glu Asn Leu Glu Lys Thr
65 70 75 80

Arg His Thr Leu Glu Lys Arg Met Asn Lys Leu Gly Thr Arg Leu Val
85 90 95

Pro Ala Glu Arg Arg Glu Lys Leu Lys Thr Ser Arg Asp Lys Leu Arg
100 105 110

Lys Ser Phe Thr Pro Asp His Val Val Tyr Ala Arg Ser Lys Thr Ala
115 120 125

Val Tyr Lys Val Pro Pro Phe Thr Phe His Val Lys Lys Ile Arg Glu
130 135 140

Gly Gln Val Glu Val Leu Lys Ala Thr Glu Met Val Glu Val Gly Ala
145 150 155 160

Asp Asp Asp Glu Gly Gly Ala Glu Arg Gly Glu Ala Gly Asp Leu Arg
165 170 175

Arg Gly Ser Ser Pro Asp Val His Ala Leu Leu Glu Ile Thr Glu Glu
180 185 190

Ser Asp Ala Val Leu Val Asp Lys Ser Asp Ser Asp
195 200

<210> 191
<211> 345
<212> PRT
<213> Homo sapiens

<400> 191

Met Ser Leu Phe Gly Leu Leu Leu Leu Thr Ser Ala Leu Ala Gly Gln
1 5 10 15

Arg Gln Gly Thr Gln Ala Glu Ser Asn Leu Ser Ser Lys Phe Gln Phe
20 25 30

Ser Ser Asn Lys Glu Gln Asn Gly Val Gln Asp Pro Gln His Glu Arg
35 40 45

Ile Ile Thr Val Ser Thr Asn Gly Ser Ile His Ser Pro Arg Phe Pro
50 55 60

His Thr Tyr Pro Arg Asn Thr Val Leu Val Trp Arg Leu Val Ala Val
 65 70 75 80

Glu Glu Asn Val Trp Ile Gln Leu Thr Phe Asp Glu Arg Phe Gly Leu
 85 90 95

Glu Asp Pro Glu Asp Asp Ile Cys Lys Tyr Asp Phe Val Glu Val Glu
 100 105 110

Glu Pro Ser Asp Gly Thr Ile Leu Gly Arg Trp Cys Gly Ser Gly Thr
 115 120 125

Val Pro Gly Lys Gln Ile Ser Lys Gly Asn Gln Ile Arg Ile Arg Phe
 130 135 140

Val Ser Asp Glu Tyr Phe Pro Ser Glu Pro Gly Phe Cys Ile His Tyr
 145 150 155 160

Asn Ile Val Met Pro Gln Phe Thr Glu Ala Val Ser Pro Ser Val Leu
 165 170 175

Pro Pro Ser Ala Leu Pro Leu Asp Leu Leu Asn Asn Ala Ile Thr Ala
 180 185 190

Phe Ser Thr Leu Glu Asp Leu Ile Arg Tyr Leu Glu Pro Glu Arg Trp
 195 200 205

Gln Leu Asp Leu Glu Asp Leu Tyr Arg Pro Thr Trp Gln Leu Leu Gly
 210 215 220

Lys Ala Phe Val Phe Gly Arg Lys Ser Arg Val Val Asp Leu Asn Leu
 225 230 235 240

Leu Thr Glu Glu Val Arg Leu Tyr Ser Cys Thr Pro Arg Asn Phe Ser
 245 250 255

Val Ser Ile Arg Glu Glu Leu Lys Arg Thr Asp Thr Ile Phe Trp Pro
 260 265 270

Gly Cys Leu Leu Val Lys Arg Cys Gly Gly Asn Cys Ala Cys Cys Leu
 275 280 285

His Asn Cys Asn Glu Cys Gln Cys Val Pro Ser Lys Val Thr Lys Lys
 290 295 300

Tyr His Glu Val Leu Gln Leu Arg Pro Lys Thr Gly Val Arg Gly Leu
 305 310 315 320

His Lys Ser Leu Thr Asp Val Ala Leu Glu His His Glu Glu Cys Asp
 325 330 335

Cys Val Cys Arg Gly Ser Thr Gly Gly
 340 345

<210> 192
 <211> 1261
 <212> PRT
 <213> Homo sapiens
 <400> 192

Met Ile Ala His Lys Gln Lys Lys Thr Lys Lys Lys Arg Ala Trp Ala
 1 5 10 15

Ser Gly Gln Leu Ser Thr Asp Ile Thr Thr Ser Glu Met Gly Leu Lys
 20 25 30

Ser Leu Ser Ser Asn Ser Ile Phe Asp Pro Asp Tyr Ile Lys Glu Leu
 35 40 45

Val Asn Asp Ile Arg Lys Phe Ser His Ile Leu Leu Tyr Leu Lys Glu
 50 55 60

Ala Ile Phe Ser Asp Cys Phe Lys Glu Val Ile His Ile Arg Leu Glu
 65 70 75 80

Glu Leu Leu Arg Val Leu Lys Ser Ile Met Asn Lys His Gln Asn Leu
 85 90 95

Asn Ser Val Asp Leu Gln Asn Ala Ala Glu Met Leu Thr Ala Lys Val
 100 105 110

Lys Ala Val Asn Phe Thr Glu Val Asn Glu Glu Asn Lys Asn Asp Leu
 115 120 125

Phe Gln Glu Val Phe Ser Ser Ile Glu Thr Leu Ala Phe Thr Phe Gly

130	135	140
Asn Ile Leu Thr Asn Phe Leu Met Gly Asp Val Gly Asn Asp Ser Phe		
145	150	155 160
Leu Arg Leu Pro Val Ser Arg Glu Thr Lys Ser Phe Glu Asn Val Ser		
	165	170 175
Val Glu Ser Val Asp Ser Ser Ser Glu Lys Gly Asn Phe Ser Pro Leu		
	180	185 190
Glu Leu Asp Asn Val Leu Leu Lys Asn Thr Asp Ser Ile Glu Leu Ala		
	195	200 205
Leu Ser Tyr Ala Lys Thr Trp Ser Lys Tyr Thr Lys Asn Ile Val Ser		
	210	215 220
Trp Val Glu Lys Lys Leu Asn Leu Glu Leu Glu Ser Thr Arg Asn Met		
	225	230 235 240
Val Lys Leu Ala Glu Ala Thr Arg Thr Asn Ile Gly Ile Gln Glu Phe		
	245	250 255
Met Pro Leu Gln Ser Leu Phe Thr Asn Ala Leu Leu Asn Asp Ile Glu		
	260	265 270
Ser Ser His Leu Leu Gln Gln Thr Ile Ala Ala Leu Gln Ala Asn Lys		
	275	280 285
Phe Val Gln Pro Leu Leu Gly Arg Lys Asn Glu Met Glu Lys Gln Arg		
	290	295 300
Lys Glu Ile Lys Glu Leu Trp Lys Gln Glu Gln Asn Lys Met Leu Glu		
	305	310 315 320
Ala Glu Asn Ala Leu Lys Lys Ala Lys Leu Leu Cys Met Gln Arg Gln		
	325	330 335
Asp Glu Tyr Glu Lys Ala Lys Ser Ser Met Phe Arg Ala Glu Glu Glu		
	340	345 350
His Leu Ser Ser Ser Gly Gly Leu Ala Lys Asn Leu Asn Lys Gln Leu		
	355	360 365

Glu Lys Lys Arg Arg Leu Glu Glu Glu Ala Leu Gln Lys Val Glu Glu
 370 375 380

Ala Asp Glu Leu Tyr Lys Val Cys Val Thr Asn Val Glu Glu Arg Arg
 385 390 395 400

Asn Asp Val Glu Asn Thr Lys Arg Glu Ile Leu Ala Gln Leu Arg Thr
 405 410 415

Leu Val Phe Gln Cys Asp Leu Thr Leu Lys Ala Val Thr Val Asn Leu
 420 425 430

Phe His Met Gln His Leu Gln Ala Ala Ser Leu Ala Asp Arg Leu Gln
 435 440 445

Ser Leu Cys Gly Ser Ala Lys Leu Tyr Asp Pro Gly Gln Glu Tyr Ser
 450 455 460

Glu Phe Val Lys Ala Thr Asn Ser Thr Glu Glu Glu Lys Val Asp Gly
 465 470 475 480

Asn Val Asn Lys His Leu Asn Ser Ser Gln Pro Ser Gly Phe Gly Pro
 485 490 495

Ala Asn Ser Leu Glu Asp Val Val Arg Leu Pro Asp Ser Ser Asn Lys
 500 505 510

Ile Glu Glu Asp Arg Cys Ser Asn Ser Ala Asp Ile Thr Gly Pro Ser
 515 520 525

Phe Ile Arg Ser Trp Thr Phe Gly Met Phe Ser Asp Ser Glu Ser Thr
 530 535 540

Gly Gly Ser Ser Glu Ser Arg Ser Leu Asp Ser Glu Ser Ile Ser Pro
 545 550 555 560

Gly Asp Phe His Arg Lys Leu Pro Arg Thr Pro Ser Ser Gly Thr Met
 565 570 575

Ser Ser Ala Asp Asp Leu Asp Glu Arg Glu Pro Pro Ser Pro Ser Glu
 580 585 590

Thr Gly Pro Asn Ser Leu Gly Thr Phe Lys Lys Thr Leu Met Ser Lys
 595 600 605

Ala Ala Leu Thr His Lys Phe Arg Lys Leu Arg Ser Pro Thr Lys Cys
 610 615 620

Arg Asp Cys Glu Gly Ile Val Val Phe Gln Gly Val Glu Cys Glu Glu
 625 630 635 640

Cys Leu Leu Val Cys His Arg Lys Cys Leu Glu Asn Leu Val Ile Ile
 645 650 655

Cys Gly His Gln Lys Leu Pro Gly Lys Ile His Leu Phe Gly Ala Glu
 660 665 670

Phe Thr Leu Val Ala Lys Lys Glu Pro Asp Gly Ile Pro Phe Ile Leu
 675 680 685

Lys Ile Cys Ala Ser Glu Ile Glu Asn Arg Ala Leu Cys Leu Gln Gly
 690 695 700

Ile Tyr Arg Val Cys Gly Asn Lys Ile Lys Thr Glu Lys Leu Cys Leu
 705 710 715 720

Ala Leu Glu Asn Gly Met His Leu Val Asp Ile Ser Glu Phe Ser Ser
 725 730 735

His Asp Ile Cys Asp Val Leu Lys Leu Tyr Leu Arg Gln Leu Pro Glu
 740 745 750

Pro Phe Ile Leu Phe Arg Leu Tyr Lys Glu Phe Ile Asp Leu Ala Lys
 755 760 765

Glu Ile Gln His Val Asn Glu Glu Gln Glu Thr Lys Lys Asn Ser Leu
 770 775 780

Glu Asp Lys Lys Trp Pro Asn Met Cys Ile Glu Ile Asn Arg Ile Leu
 785 790 795 800

Leu Lys Ser Lys Asp Leu Leu Arg Gln Leu Pro Ala Ser Asn Phe Asn
 805 810 815

Ser Leu His Phe Leu Ile Val His Leu Lys Arg Val Val Asp His Ala
 820 825 830
 Glu Glu Asn Lys Met Asn Ser Lys Asn Leu Gly Val Ile Phe Gly Pro
 835 840 845
 Ser Leu Ile Arg Pro Arg Pro Gln Thr Ala Pro Ile Thr Ile Ser Ser
 850 855 860
 Leu Ala Glu Tyr Ser Asn Gln Ala Arg Leu Val Glu Phe Leu Ile Thr
 865 870 875 880
 Tyr Ser Gln Lys Ile Phe Asp Gly Ser Leu Gln Pro Gln Asp Val Met
 885 890 895
 Cys Ser Ile Gly Val Val Asp Gln Gly Cys Phe Pro Lys Pro Leu Leu
 900 905 910
 Ser Pro Glu Glu Arg Asp Ile Glu Arg Ser Met Lys Ser Leu Phe Phe
 915 920 925
 Ser Ser Lys Glu Asp Ile His Thr Ser Glu Ser Glu Ser Lys Ile Phe
 930 935 940
 Glu Arg Ala Thr Ser Phe Glu Glu Ser Glu Arg Lys Gln Asn Ala Leu
 945 950 955 960
 Gly Lys Cys Asp Ala Cys Leu Ser Asp Lys Ala Gln Leu Leu Leu Asp
 965 970 975
 Gln Glu Ala Glu Ser Ala Ser Gln Lys Ile Glu Asp Gly Lys Ala Pro
 980 985 990
 Lys Pro Leu Ser Leu Lys Ser Asp Arg Ser Thr Asn Asn Val Glu Arg
 995 1000 1005
 His Thr Pro Arg Thr Lys Ile Arg Pro Val Ser Leu Pro Val Asp
 1010 1015 1020
 Arg Leu Leu Leu Ala Ser Pro Pro Asn Glu Arg Asn Gly Arg Asn
 1025 1030 1035
 Met Gly Asn Val Asn Leu Asp Lys Phe Cys Lys Asn Pro Ala Phe

1040		1045		1050
Glu Gly Val Asn Arg Lys Asp	Ala Ala Thr Thr Val	Cys Ser Lys		
1055	1060	1065		
Phe Asn Gly Phe Asp Gln Gln	Thr Leu Gln Lys Ile	Gln Asp Lys		
1070	1075	1080		
Gln Tyr Glu Gln Asn Ser Leu	Thr Ala Lys Thr Thr	Met Ile Met		
1085	1090	1095		
Pro Ser Ala Leu Gln Glu Lys	Gly Val Thr Thr Ser	Leu Gln Ile		
1100	1105	1110		
Ser Gly Asp His Ser Ile Asn	Ala Thr Gln Pro Ser	Lys Pro Tyr		
1115	1120	1125		
Ala Glu Pro Val Arg Ser Val	Arg Glu Ala Ser Glu	Arg Arg Ser		
1130	1135	1140		
Ser Asp Ser Tyr Pro Leu Ala	Pro Val Arg Ala Pro	Arg Thr Leu		
1145	1150	1155		
Gln Pro Gln His Trp Thr Thr	Phe Tyr Lys Pro His	Ala Pro Ile		
1160	1165	1170		
Ile Ser Ile Arg Gly Asn Glu	Glu Lys Pro Ala Ser	Pro Ser Ala		
1175	1180	1185		
Ala Cys Pro Pro Gly Thr Asp	His Asp Pro His Gly	Leu Val Val		
1190	1195	1200		
Lys Ser Met Pro Asp Pro Asp	Lys Ala Ser Ala Cys	Pro Gly Gln		
1205	1210	1215		
Ala Thr Gly Gln Pro Lys Glu	Asp Ser Glu Glu Leu	Gly Leu Pro		
1220	1225	1230		
Asp Val Asn Pro Met Cys Gln	Arg Pro Arg Leu Lys	Arg Met Gln		
1235	1240	1245		
Gln Phe Glu Asp Leu Glu Asp	Glu Ile Pro Gln Phe	Val		
1250	1255	1260		

<210> 193
 <211> 192
 <212> PRT
 <213> Homo sapiens

<400> 193

Met Gln Ala Ile Lys Cys Val Val Val Gly Asp Gly Ala Val Gly Lys
 1 5 10 15

Thr Cys Leu Leu Ile Ser Tyr Thr Thr Asn Ala Phe Pro Gly Glu Tyr
 20 25 30

Ile Pro Thr Val Phe Asp Asn Tyr Ser Ala Asn Val Met Val Asp Ser
 35 40 45

Lys Pro Val Asn Leu Gly Leu Trp Asp Thr Ala Gly Gln Glu Asp Tyr
 50 55 60

Asp Arg Leu Arg Pro Leu Ser Tyr Pro Gln Thr Asp Val Phe Leu Ile
 65 70 75 80

Cys Phe Ser Leu Val Ser Pro Ala Ser Tyr Glu Asn Val Arg Ala Lys
 85 90 95

Trp Phe Pro Glu Val Arg His His Cys Pro Ser Thr Pro Ile Ile Leu
 100 105 110

Val Gly Thr Lys Leu Asp Leu Arg Asp Asp Lys Asp Thr Ile Glu Lys
 115 120 125

Leu Lys Glu Lys Lys Leu Ala Pro Ile Thr Tyr Pro Gln Gly Leu Ala
 130 135 140

Leu Ala Lys Glu Ile Asp Ser Val Lys Tyr Leu Glu Cys Ser Ala Leu
 145 150 155 160

Thr Gln Arg Gly Leu Lys Thr Val Phe Asp Glu Ala Ile Arg Ala Val
 165 170 175

Leu Cys Pro Gln Pro Thr Arg Gln Gln Lys Arg Ala Cys Ser Leu Leu
 180 185 190

<210> 194
 <211> 404
 <212> PRT
 <213> Homo sapiens

<400> 194

Met Asp Ser Arg Thr Lys Ser Lys Asp Tyr Cys Lys Val Ile Phe Pro
 1 5 10 15

Tyr Glu Ala Gln Asn Asp Asp Glu Leu Thr Ile Lys Glu Gly Asp Ile
 20 25 30

Val Thr Leu Ile Asn Lys Asp Cys Ile Asp Val Gly Trp Trp Glu Gly
 35 40 45

Glu Leu Asn Gly Arg Arg Gly Val Phe Pro Asp Asn Phe Val Lys Leu
 50 55 60

Leu Pro Pro Asp Phe Glu Lys Glu Gly Asn Arg Pro Lys Lys Pro Pro
 65 70 75 80

Pro Pro Ser Ala Pro Val Ile Lys Gln Gly Ala Gly Thr Thr Glu Arg
 85 90 95

Lys His Glu Ile Lys Lys Ile Pro Pro Glu Arg Pro Glu Met Leu Pro
 100 105 110

Asn Arg Thr Glu Glu Lys Glu Arg Pro Glu Arg Glu Pro Lys Leu Asp
 115 120 125

Leu Gln Lys Pro Ser Val Pro Ala Ile Pro Pro Lys Lys Pro Arg Pro
 130 135 140

Pro Lys Thr Asn Ser Leu Ser Arg Pro Gly Ala Leu Pro Pro Arg Arg
 145 150 155 160

Pro Glu Arg Pro Val Gly Pro Leu Thr His Thr Arg Gly Asp Ser Pro
 165 170 175

Lys Ile Asp Leu Ala Gly Ser Ser Leu Ser Gly Ile Leu Asp Lys Asp
 180 185 190

Leu Ser Asp Arg Ser Asn Asp Ile Asp Leu Glu Gly Phe Asp Ser Val
 195 200 205

Val Ser Ser Thr Glu Lys Leu Ser His Pro Thr Thr Ser Arg Pro Lys
 210 215 220

Ala Thr Gly Arg Arg Pro Pro Ser Gln Ser Leu Thr Ser Ser Ser Leu
 225 230 235 240

Ser Ser Pro Asp Ile Phe Asp Ser Pro Ser Pro Glu Glu Asp Lys Glu
 245 250 255

Glu His Ile Ser Leu Ala His Arg Gly Val Asp Ala Ser Lys Lys Thr
 260 265 270

Ser Lys Thr Val Thr Ile Ser Gln Val Ser Asp Asn Lys Ala Ser Leu
 275 280 285

Pro Pro Lys Pro Gly Thr Met Ala Ala Gly Gly Gly Gly Pro Ala Pro
 290 295 300

Leu Ser Ser Ala Ala Pro Ser Pro Leu Ser Ser Ser Leu Gly Thr Ala
 305 310 315 320

Gly His Arg Ala Asn Ser Pro Ser Leu Phe Gly Thr Glu Gly Lys Pro
 325 330 335

Lys Met Glu Pro Ala Ala Ser Ser Gln Ala Ala Val Glu Glu Leu Arg
 340 345 350

Thr Gln Val Arg Glu Leu Arg Ser Ile Ile Glu Thr Met Lys Asp Gln
 355 360 365

Gln Lys Arg Glu Ile Lys Gln Leu Leu Ser Glu Leu Asp Glu Glu Lys
 370 375 380

Lys Ile Arg Leu Arg Leu Gln Met Glu Val Asn Asp Ile Lys Lys Ala
 385 390 395 400

Leu Gln Ser Lys

<210> 195
 <211> 268
 <212> PRT

<213> Homo sapiens

<400> 195

Met Pro Arg Ser Phe Leu Val Lys Lys His Phe Asn Ala Ser Lys Lys
 1 5 10 15

Pro Asn Tyr Ser Glu Leu Asp Thr His Thr Val Ile Ile Ser Pro Tyr
 20 25 30

Leu Tyr Glu Ser Tyr Ser Met Pro Val Ile Pro Gln Pro Glu Ile Leu
 35 40 45

Ser Ser Gly Ala Tyr Ser Pro Ile Thr Val Trp Thr Thr Ala Ala Pro
 50 55 60

Phe His Ala Gln Leu Pro Asn Gly Leu Ser Pro Leu Ser Gly Tyr Ser
 65 70 75 80

Ser Ser Leu Gly Arg Val Ser Pro Pro Pro Pro Ser Asp Thr Ser Ser
 85 90 95

Lys Asp His Ser Gly Ser Glu Ser Pro Ile Ser Asp Glu Glu Glu Arg
 100 105 110

Leu Gln Ser Lys Leu Ser Asp Pro His Ala Ile Glu Ala Glu Lys Phe
 115 120 125

Gln Cys Asn Leu Cys Asn Lys Thr Tyr Ser Thr Phe Ser Gly Leu Ala
 130 135 140

Lys His Lys Gln Leu His Cys Asp Ala Gln Ser Arg Lys Ser Phe Ser
 145 150 155 160

Cys Lys Tyr Cys Asp Lys Glu Tyr Val Ser Leu Gly Ala Leu Lys Met
 165 170 175

His Ile Arg Thr His Thr Leu Pro Cys Val Cys Lys Ile Cys Gly Lys
 180 185 190

Ala Phe Ser Arg Pro Trp Leu Leu Gln Gly His Ile Arg Thr His Thr
 195 200 205

Gly Glu Lys Pro Phe Ser Cys Pro His Cys Asn Arg Ala Phe Ala Asp

210 215 220
 Arg Ser Asn Leu Arg Ala His Leu Gln Thr His Ser Asp Val Lys Lys
 225 230 235 240
 Tyr Gln Cys Lys Asn Cys Ser Lys Thr Phe Ser Arg Met Ser Leu Leu
 245 250 255
 His Lys His Glu Glu Ser Gly Cys Cys Val Ala His
 260 265

 <210> 196
 <211> 490
 <212> PRT
 <213> Homo sapiens

 <400> 196
 Met Ser Glu Val Thr Lys Asn Ser Leu Glu Lys Ile Leu Pro Gln Leu
 1 5 10 15
 Lys Cys His Phe Thr Trp Asn Leu Phe Lys Glu Asp Ser Val Ser Arg
 20 25 30
 Asp Leu Glu Asp Arg Val Cys Asn Gln Ile Glu Phe Leu Asn Thr Glu
 35 40 45
 Phe Lys Ala Thr Met Tyr Asn Leu Leu Ala Tyr Ile Lys His Leu Asp
 50 55 60
 Gly Asn Asn Glu Ala Ala Leu Glu Cys Leu Arg Gln Ala Glu Glu Leu
 65 70 75 80
 Ile Gln Gln Glu His Ala Asp Gln Ala Glu Ile Arg Ser Leu Val Thr
 85 90 95
 Trp Gly Asn Tyr Ala Trp Val Tyr Tyr His Leu Gly Arg Leu Ser Asp
 100 105 110
 Ala Gln Ile Tyr Val Asp Lys Val Lys Gln Thr Cys Lys Lys Phe Ser
 115 120 125
 Asn Pro Tyr Ser Ile Glu Tyr Ser Glu Leu Asp Cys Glu Glu Gly Trp
 130 135 140

Thr Gln Leu Lys Cys Gly Arg Asn Glu Arg Ala Lys Val Cys Phe Glu
145 150 155 160

Lys Ala Leu Glu Glu Lys Pro Asn Asn Pro Glu Phe Ser Ser Gly Leu
165 170 175

Ala Ile Ala Met Tyr His Leu Asp Asn His Pro Glu Lys Gln Phe Ser
180 185 190

Thr Asp Val Leu Lys Gln Ala Ile Glu Leu Ser Pro Asp Asn Gln Tyr
195 200 205

Val Lys Val Leu Leu Gly Leu Lys Leu Gln Lys Met Asn Lys Glu Ala
210 215 220

Glu Gly Glu Gln Phe Val Glu Glu Ala Leu Glu Lys Ser Pro Cys Gln
225 230 235 240

Thr Asp Val Leu Arg Ser Ala Ala Lys Phe Tyr Arg Arg Lys Gly Asp
245 250 255

Leu Asp Lys Ala Ile Glu Leu Phe Gln Arg Val Leu Glu Ser Thr Pro
260 265 270

Asn Asn Gly Tyr Leu Tyr His Gln Ile Gly Cys Cys Tyr Lys Ala Lys
275 280 285

Val Arg Gln Met Gln Asn Thr Gly Glu Ser Glu Ala Ser Gly Asn Lys
290 295 300

Glu Met Ile Glu Ala Leu Lys Gln Tyr Ala Met Asp Tyr Ser Asn Lys
305 310 315 320

Ala Leu Glu Lys Gly Leu Asn Pro Leu Asn Ala Tyr Ser Asp Leu Ala
325 330 335

Glu Phe Leu Glu Thr Glu Cys Tyr Gln Thr Pro Phe Asn Lys Glu Val
340 345 350

Pro Asp Ala Glu Lys Gln Gln Ser His Gln Arg Tyr Cys Asn Leu Gln
355 360 365

Lys Tyr Asn Gly Lys Ser Glu Asp Thr Ala Val Gln His Gly Leu Glu
 370 375 380

Gly Leu Ser Ile Ser Lys Lys Ser Thr Asp Lys Glu Glu Ile Lys Asp
 385 390 395 400

Gln Pro Gln Asn Val Ser Glu Asn Leu Leu Pro Gln Asn Ala Pro Asn
 405 410 415

Tyr Trp Tyr Leu Gln Gly Leu Ile His Lys Gln Asn Gly Asp Leu Leu
 420 425 430

Gln Ala Ala Lys Cys Tyr Glu Lys Glu Leu Gly Arg Leu Leu Arg Asp
 435 440 445

Ala Pro Ser Gly Ile Gly Ser Ile Phe Leu Ser Ala Ser Glu Leu Glu
 450 455 460

Asp Gly Ser Glu Glu Met Gly Gln Gly Ala Val Ser Ser Ser Pro Arg
 465 470 475 480

Glu Leu Leu Ser Asn Ser Glu Gln Leu Asn
 485 490

<210> 197
 <211> 567
 <212> PRT
 <213> Homo sapiens

<400> 197

Met Gly Arg Gly Leu Leu Arg Gly Leu Trp Pro Leu His Ile Val Leu
 1 5 10 15

Trp Thr Arg Ile Ala Ser Thr Ile Pro Pro His Val Gln Lys Ser Val
 20 25 30

Asn Asn Asp Met Ile Val Thr Asp Asn Asn Gly Ala Val Lys Phe Pro
 35 40 45

Gln Leu Cys Lys Phe Cys Asp Val Arg Phe Ser Thr Cys Asp Asn Gln
 50 55 60

Lys Ser Cys Met Ser Asn Cys Ser Ile Thr Ser Ile Cys Glu Lys Pro
 65 70 75 80

Gln Glu Val Cys Val Ala Val Trp Arg Lys Asn Asp Glu Asn Ile Thr
85 90 95

Leu Glu Thr Val Cys His Asp Pro Lys Leu Pro Tyr His Asp Phe Ile
100 105 110

Leu Glu Asp Ala Ala Ser Pro Lys Cys Ile Met Lys Glu Lys Lys Lys
115 120 125

Pro Gly Glu Thr Phe Phe Met Cys Ser Cys Ser Ser Asp Glu Cys Asn
130 135 140

Asp Asn Ile Ile Phe Ser Glu Glu Tyr Asn Thr Ser Asn Pro Asp Leu
145 150 155 160

Leu Leu Val Ile Phe Gln Val Thr Gly Ile Ser Leu Leu Pro Pro Leu
165 170 175

Gly Val Ala Ile Ser Val Ile Ile Ile Phe Tyr Cys Tyr Arg Val Asn
180 185 190

Arg Gln Gln Lys Leu Ser Ser Thr Trp Glu Thr Gly Lys Thr Arg Lys
195 200 205

Leu Met Glu Phe Ser Glu His Cys Ala Ile Ile Leu Glu Asp Asp Arg
210 215 220

Ser Asp Ile Ser Ser Thr Cys Ala Asn Asn Ile Asn His Asn Thr Glu
225 230 235 240

Leu Leu Pro Ile Glu Leu Asp Thr Leu Val Gly Lys Gly Arg Phe Ala
245 250 255

Glu Val Tyr Lys Ala Lys Leu Lys Gln Asn Thr Ser Glu Gln Phe Glu
260 265 270

Thr Val Ala Val Lys Ile Phe Pro Tyr Glu Glu Tyr Ala Ser Trp Lys
275 280 285

Thr Glu Lys Asp Ile Phe Ser Asp Ile Asn Leu Lys His Glu Asn Ile
290 295 300

Leu Gln Phe Leu Thr Ala Glu Glu Arg Lys Thr Glu Leu Gly Lys Gln
 305 310 315 320

Tyr Trp Leu Ile Thr Ala Phe His Ala Lys Gly Asn Leu Gln Glu Tyr
 325 330 335

Leu Thr Arg His Val Ile Ser Trp Glu Asp Leu Arg Lys Leu Gly Ser
 340 345 350

Ser Leu Ala Arg Gly Ile Ala His Leu His Ser Asp His Thr Pro Cys
 355 360 365

Gly Arg Pro Lys Met Pro Ile Val His Arg Asp Leu Asn Ser Ser Asn
 370 375 380

Ile Leu Val Lys Asn Asp Leu Thr Cys Cys Leu Cys Asp Phe Gly Leu
 385 390 395 400

Ser Leu Arg Leu Asp Pro Thr Leu Ser Val Asp Asp Leu Ala Asn Ser
 405 410 415

Gly Gln Val Gly Thr Ala Arg Tyr Met Ala Pro Glu Val Leu Glu Ser
 420 425 430

Arg Met Asn Leu Glu Asn Ala Glu Ser Phe Lys Gln Thr Asp Val Tyr
 435 440 445

Ser Met Ala Leu Val Leu Trp Glu Met Thr Ser Arg Cys Asn Ala Val
 450 455 460

Gly Glu Val Lys Asp Tyr Glu Pro Pro Phe Gly Ser Lys Val Arg Glu
 465 470 475 480

His Pro Cys Val Glu Ser Met Lys Asp Asn Val Leu Arg Asp Arg Gly
 485 490 495

Arg Pro Glu Ile Pro Ser Phe Trp Leu Asn His Gln Gly Ile Gln Met
 500 505 510

Val Cys Glu Thr Leu Thr Glu Cys Trp Asp His Asp Pro Glu Ala Arg
 515 520 525

Leu Thr Ala Gln Cys Val Ala Glu Arg Phe Ser Glu Leu Glu His Leu
 530 535 540

Asp Arg Leu Ser Gly Arg Ser Cys Ser Glu Glu Lys Ile Pro Glu Asp
 545 550 555 560

Gly Ser Leu Asn Thr Thr Lys
 565

<210> 198
 <211> 425
 <212> PRT
 <213> Homo sapiens

<400> 198

Met Ser Ser Ile Leu Pro Phe Thr Pro Pro Ile Val Lys Arg Leu Leu
 1 5 10 15

Gly Trp Lys Lys Gly Glu Gln Asn Gly Gln Glu Glu Lys Trp Cys Glu
 20 25 30

Lys Ala Val Lys Ser Leu Val Lys Lys Leu Lys Lys Thr Gly Gln Leu
 35 40 45

Asp Glu Leu Glu Lys Ala Ile Thr Thr Gln Asn Val Asn Thr Lys Cys
 50 55 60

Ile Thr Ile Pro Arg Ser Leu Asp Gly Arg Leu Gln Val Ser His Arg
 65 70 75 80

Lys Gly Leu Pro His Val Ile Tyr Cys Arg Leu Trp Arg Trp Pro Asp
 85 90 95

Leu His Ser His His Glu Leu Arg Ala Met Glu Leu Cys Glu Phe Ala
 100 105 110

Phe Asn Met Lys Lys Asp Glu Val Cys Val Asn Pro Tyr His Tyr Gln
 115 120 125

Arg Val Glu Thr Pro Val Leu Pro Pro Val Leu Val Pro Arg His Thr
 130 135 140

Glu Ile Pro Ala Glu Phe Pro Pro Leu Asp Asp Tyr Ser His Ser Ile
 145 150 155 160

Pro Glu Asn Thr Asn Phe Pro Ala Gly Ile Glu Pro Gln Ser Asn Ile
 165 170 175

Pro Glu Thr Pro Pro Pro Gly Tyr Leu Ser Glu Asp Gly Glu Thr Ser
 180 185 190

Asp His Gln Met Asn His Ser Met Asp Ala Gly Ser Pro Asn Leu Ser
 195 200 205

Pro Asn Pro Met Ser Pro Ala His Asn Asn Leu Asp Leu Gln Pro Val
 210 215 220

Thr Tyr Cys Glu Pro Ala Phe Trp Cys Ser Ile Ser Tyr Tyr Glu Leu
 225 230 235 240

Asn Gln Arg Val Gly Glu Thr Phe His Ala Ser Gln Pro Ser Met Thr
 245 250 255

Val Asp Gly Phe Thr Asp Pro Ser Asn Ser Glu Arg Phe Cys Leu Gly
 260 265 270

Leu Leu Ser Asn Val Asn Arg Asn Ala Ala Val Glu Leu Thr Arg Arg
 275 280 285

His Ile Gly Arg Gly Val Arg Leu Tyr Tyr Ile Gly Gly Glu Val Phe
 290 295 300

Ala Glu Cys Leu Ser Asp Ser Ala Ile Phe Val Gln Ser Pro Asn Cys
 305 310 315 320

Asn Gln Arg Tyr Gly Trp His Pro Ala Thr Val Cys Lys Ile Pro Pro
 325 330 335

Gly Cys Asn Leu Lys Ile Phe Asn Asn Gln Glu Phe Ala Ala Leu Leu
 340 345 350

Ala Gln Ser Val Asn Gln Gly Phe Glu Ala Val Tyr Gln Leu Thr Arg
 355 360 365

Met Cys Thr Ile Arg Met Ser Phe Val Lys Gly Trp Gly Ala Glu Tyr
 370 375 380

Arg Arg Gln Thr Val Thr Ser Thr Pro Cys Trp Ile Glu Leu His Leu
385 390 395 400

Asn Gly Pro Leu Gln Trp Leu Asp Lys Val Leu Thr Gln Met Gly Ser
405 410 415

Pro Ser Ile Arg Cys Ser Ser Val Ser
420 425

<210> 199
<211> 655
<212> PRT
<213> Homo sapiens

<400> 199

Met Gly Thr Ser Pro Ser Ser Ser Thr Ala Leu Ala Ser Cys Ser Arg
1 5 10 15

Ile Ala Arg Arg Ala Thr Ala Thr Met Ile Ala Gly Ser Leu Leu Leu
20 25 30

Leu Gly Phe Leu Ser Thr Thr Thr Ala Gln Pro Glu Gln Lys Ala Ser
35 40 45

Asn Leu Ile Gly Thr Tyr Arg His Val Asp Arg Ala Thr Gly Gln Val
50 55 60

Leu Thr Cys Asp Lys Cys Pro Ala Gly Thr Tyr Val Ser Glu His Cys
65 70 75 80

Thr Asn Thr Ser Leu Arg Val Cys Ser Ser Cys Pro Val Gly Thr Phe
85 90 95

Thr Arg His Glu Asn Gly Ile Glu Lys Cys His Asp Cys Ser Gln Pro
100 105 110

Cys Pro Trp Pro Met Ile Glu Lys Leu Pro Cys Ala Ala Leu Thr Asp
115 120 125

Arg Glu Cys Thr Cys Pro Pro Gly Met Phe Gln Ser Asn Ala Thr Cys
130 135 140

Ala Pro His Thr Val Cys Pro Val Gly Trp Gly Val Arg Lys Lys Gly

145 150 155 160
 Thr Glu Thr Glu Asp Val Arg Cys Lys Gln Cys Ala Arg Gly Thr Phe
 165 170 175
 Ser Asp Val Pro Ser Ser Val Met Lys Cys Lys Ala Tyr Thr Asp Cys
 180 185 190
 Leu Ser Gln Asn Leu Val Val Ile Lys Pro Gly Thr Lys Glu Thr Asp
 195 200 205
 Asn Val Cys Gly Thr Leu Pro Ser Phe Ser Ser Ser Thr Ser Pro Ser
 210 215 220
 Pro Gly Thr Ala Ile Phe Pro Arg Pro Glu His Met Glu Thr His Glu
 225 230 235 240
 Val Pro Ser Ser Thr Tyr Val Pro Lys Gly Met Asn Ser Thr Glu Ser
 245 250 255
 Asn Ser Ser Ala Ser Val Arg Pro Lys Val Leu Ser Ser Ile Gln Glu
 260 265 270
 Gly Thr Val Pro Asp Asn Thr Ser Ser Ala Arg Gly Lys Glu Asp Val
 275 280 285
 Asn Lys Thr Leu Pro Asn Leu Gln Val Val Asn His Gln Gln Gly Pro
 290 295 300
 His His Arg His Ile Leu Lys Leu Leu Pro Ser Met Glu Ala Thr Gly
 305 310 315 320
 Gly Glu Lys Ser Ser Thr Pro Ile Lys Gly Pro Lys Arg Gly His Pro
 325 330 335
 Arg Gln Asn Leu His Lys His Phe Asp Ile Asn Glu His Leu Pro Trp
 340 345 350
 Met Ile Val Leu Phe Leu Leu Leu Val Leu Val Val Ile Val Val Cys
 355 360 365
 Ser Ile Arg Lys Ser Ser Arg Thr Leu Lys Lys Gly Pro Arg Gln Asp
 370 375 380

Pro Ser Ala Ile Val Glu Lys Ala Gly Leu Lys Lys Ser Met Thr Pro
 385 390 395 400

Thr Gln Asn Arg Glu Lys Trp Ile Tyr Tyr Cys Asn Gly His Gly Ile
 405 410 415

Asp Ile Leu Lys Leu Val Ala Ala Gln Val Gly Ser Gln Trp Lys Asp
 420 425 430

Ile Tyr Gln Phe Leu Cys Asn Ala Ser Glu Arg Glu Val Ala Ala Phe
 435 440 445

Ser Asn Gly Tyr Thr Ala Asp His Glu Arg Ala Tyr Ala Ala Leu Gln
 450 455 460

His Trp Thr Ile Arg Gly Pro Glu Ala Ser Leu Ala Gln Leu Ile Ser
 465 470 475 480

Ala Leu Arg Gln His Arg Arg Asn Asp Val Val Glu Lys Ile Arg Gly
 485 490 495

Leu Met Glu Asp Thr Thr Gln Leu Glu Thr Asp Lys Leu Ala Leu Pro
 500 505 510

Met Ser Pro Ser Pro Leu Ser Pro Ser Pro Ile Pro Ser Pro Asn Ala
 515 520 525

Lys Leu Glu Asn Ser Ala Leu Leu Thr Val Glu Pro Ser Pro Gln Asp
 530 535 540

Lys Asn Lys Gly Phe Phe Val Asp Glu Ser Glu Pro Leu Leu Arg Cys
 545 550 555 560

Asp Ser Thr Ser Ser Gly Ser Ser Ala Leu Ser Arg Asn Gly Ser Phe
 565 570 575

Ile Thr Lys Glu Lys Lys Asp Thr Val Leu Arg Gln Val Arg Leu Asp
 580 585 590

Pro Cys Asp Leu Gln Pro Ile Phe Asp Asp Met Leu His Phe Leu Asn
 595 600 605

Pro Glu Glu Leu Arg Val Ile Glu Glu Ile Pro Gln Ala Glu Asp Lys
 610 615 620

Leu Asp Arg Leu Phe Glu Ile Ile Gly Val Lys Ser Gln Glu Ala Ser
 625 630 635 640

Gln Thr Leu Leu Asp Ser Val Tyr Ser His Leu Pro Asp Leu Leu
 645 650 655

<210> 200
 <211> 207
 <212> PRT
 <213> Homo sapiens

<400> 200

Met Ser Ser Asp Arg Gln Arg Ser Asp Asp Glu Ser Pro Ser Thr Ser
 1 5 10 15

Ser Gly Ser Ser Asp Ala Asp Gln Arg Asp Pro Ala Ala Pro Glu Pro
 20 25 30

Glu Glu Gln Glu Glu Arg Lys Pro Ser Ala Thr Gln Gln Lys Lys Asn
 35 40 45

Thr Lys Leu Ser Ser Lys Thr Thr Ala Lys Leu Ser Thr Ser Ala Lys
 50 55 60

Arg Ile Gln Lys Glu Leu Ala Glu Ile Thr Leu Asp Pro Pro Pro Asn
 65 70 75 80

Cys Ser Ala Gly Pro Lys Gly Asp Asn Ile Tyr Glu Trp Arg Ser Thr
 85 90 95

Ile Leu Gly Pro Pro Gly Ser Val Tyr Glu Gly Gly Val Phe Phe Leu
 100 105 110

Asp Ile Thr Phe Ser Ser Asp Tyr Pro Phe Lys Pro Pro Lys Val Thr
 115 120 125

Phe Arg Thr Arg Ile Tyr His Cys Asn Ile Asn Ser Gln Gly Val Ile
 130 135 140

Cys Leu Asp Ile Leu Lys Asp Asn Trp Ser Pro Ala Leu Thr Ile Ser

145 150 155 160
 Lys Val Leu Leu Ser Ile Cys Ser Leu Leu Thr Asp Cys Asn Pro Ala
 165 170 175
 Asp Pro Leu Val Gly Ser Ile Ala Thr Gln Tyr Leu Thr Asn Arg Ala
 180 185 190
 Glu His Asp Arg Ile Ala Arg Gln Trp Thr Lys Arg Tyr Ala Thr
 195 200 205

 <210> 201
 <211> 572
 <212> PRT
 <213> Homo sapiens

 <400> 201
 Met Ala Ala Pro Arg Pro Ser Pro Ala Ile Ser Val Ser Val Ser Ala
 1 5 10 15

 Pro Ala Phe Tyr Ala Pro Gln Lys Lys Phe Gly Pro Val Val Ala Pro
 20 25 30

 Lys Pro Lys Val Asn Pro Phe Arg Pro Gly Asp Ser Glu Pro Pro Pro
 35 40 45

 Ala Pro Gly Ala Gln Arg Ala Gln Met Gly Arg Val Gly Glu Ile Pro
 50 55 60

 Pro Pro Pro Pro Glu Asp Phe Pro Leu Pro Pro Pro Pro Leu Ala Gly
 65 70 75 80

 Asp Gly Asp Asp Ala Glu Gly Ala Leu Gly Gly Ala Phe Pro Pro Pro
 85 90 95

 Pro Pro Pro Ile Glu Glu Ser Phe Pro Pro Ala Pro Leu Glu Glu Glu
 100 105 110

 Ile Phe Pro Ser Pro Pro Pro Pro Pro Glu Glu Glu Gly Gly Pro Glu
 115 120 125

 Ala Pro Ile Pro Pro Pro Pro Gln Pro Arg Glu Lys Val Ser Ser Ile
 130 135 140

Asp Leu Glu Ile Asp Ser Leu Ser Ser Leu Leu Asp Asp Met Thr Lys
 145 150 155 160

Asn Asp Pro Phe Lys Ala Arg Val Ser Ser Gly Tyr Val Pro Pro Pro
 165 170 175

Val Ala Thr Pro Phe Ser Ser Lys Ser Ser Thr Lys Pro Ala Ala Gly
 180 185 190

Gly Thr Ala Pro Leu Pro Pro Trp Lys Ser Pro Ser Ser Ser Gln Pro
 195 200 205

Leu Pro Gln Val Pro Ala Pro Ala Gln Ser Gln Thr Gln Phe His Val
 210 215 220

Gln Pro Gln Pro Gln Pro Lys Pro Gln Val Gln Leu His Val Gln Ser
 225 230 235 240

Gln Thr Gln Pro Val Ser Leu Ala Asn Thr Gln Pro Arg Gly Pro Pro
 245 250 255

Ala Ser Ser Pro Ala Pro Ala Pro Lys Phe Ser Pro Val Thr Pro Lys
 260 265 270

Phe Thr Pro Val Ala Ser Lys Phe Ser Pro Gly Ala Pro Gly Gly Ser
 275 280 285

Gly Ser Gln Pro Asn Gln Lys Leu Gly His Pro Glu Ala Leu Ser Ala
 290 295 300

Gly Thr Gly Ser Pro Gln Pro Pro Ser Phe Thr Tyr Ala Gln Gln Arg
 305 310 315 320

Glu Lys Pro Arg Val Gln Glu Lys Gln His Pro Val Pro Pro Pro Ala
 325 330 335

Gln Asn Gln Asn Gln Val Arg Ser Pro Gly Ala Pro Gly Pro Leu Thr
 340 345 350

Leu Lys Glu Val Glu Glu Leu Glu Gln Leu Thr Gln Gln Leu Met Gln
 355 360 365

Asp Met Glu His Pro Gln Arg Gln Asn Val Ala Val Asn Glu Leu Cys
 370 375 380

Gly Arg Cys His Gln Pro Leu Ala Arg Ala Gln Pro Ala Val Arg Ala
 385 390 395 400

Leu Gly Gln Leu Phe His Ile Ala Cys Phe Thr Cys His Gln Cys Ala
 405 410 415

Gln Gln Leu Gln Gly Gln Gln Phe Tyr Ser Leu Glu Gly Ala Pro Tyr
 420 425 430

Cys Glu Gly Cys Tyr Thr Asp Thr Leu Glu Lys Cys Asn Thr Cys Gly
 435 440 445

Glu Pro Ile Thr Asp Arg Met Leu Arg Ala Thr Gly Lys Ala Tyr His
 450 455 460

Pro His Cys Phe Thr Cys Val Val Cys Ala Arg Pro Leu Glu Gly Thr
 465 470 475 480

Ser Phe Ile Val Asp Gln Ala Asn Arg Pro His Cys Val Pro Asp Tyr
 485 490 495

His Lys Gln Tyr Ala Pro Arg Cys Ser Val Cys Ser Glu Pro Ile Met
 500 505 510

Pro Glu Pro Gly Arg Asp Glu Thr Val Arg Val Val Ala Leu Asp Lys
 515 520 525

Asn Phe His Met Lys Cys Tyr Lys Cys Glu Asp Cys Gly Lys Pro Leu
 530 535 540

Ser Ile Glu Ala Asp Asp Asn Gly Cys Phe Pro Leu Asp Gly His Val
 545 550 555 560

Leu Cys Arg Lys Cys His Thr Ala Arg Ala Gln Thr
 565 570

<210> 202
 <211> 141
 <212> PRT
 <213> Homo sapiens

<400> 202

Met Thr Lys Gln His Glu Leu Gly Gly Leu Leu Ala Leu Val Gln Asn
 1 5 10 15

Cys Gln Ser Glu Met Asn Ile Lys Asp Ser Arg Ala Val Gly Leu Ser
 20 25 30

Val Lys Arg Leu Cys Ile Ser Phe Val Asp Glu Phe Cys Glu Arg Thr
 35 40 45

Glu Arg Pro Leu Tyr Leu Ala Gln Gly Leu Phe Met Lys Arg Glu Thr
 50 55 60

Tyr Trp Glu Val Gln Asp Ser Gly Ile Ser Pro Leu Leu Leu Leu Leu
 65 70 75 80

Ser Thr Ala Leu Asp Cys Ser Pro Glu Ala Glu Thr Arg Gln Ser Pro
 85 90 95

Gly Gly Arg Lys Met Leu Gln Glu Pro Thr Leu Ser Met Ser Leu Gln
 100 105 110

Ile Leu Thr Gly Phe Leu Trp Val Gln Leu Trp Asn Trp Glu Thr Phe
 115 120 125

Leu Arg Ile Arg Thr His Ser Thr Asp Ala Ser Cys Pro
 130 135 140

<210> 203

<211> 430

<212> PRT

<213> Homo sapiens

<400> 203

Met Asp Glu Gln Pro Arg Leu Met His Ser His Ala Gly Val Gly Met
 1 5 10 15

Ala Gly His Pro Gly Leu Ser Gln His Leu Gln Asp Gly Ala Gly Gly
 20 25 30

Thr Glu Gly Glu Gly Gly Arg Lys Gln Asp Ile Gly Asp Ile Leu Gln
 35 40 45

Gln Ile Met Thr Ile Thr Asp Gln Ser Leu Asp Glu Ala Gln Ala Arg
 50 55 60

Lys His Ala Leu Asn Cys His Arg Met Lys Pro Ala Leu Phe Asn Val
 65 70 75 80

Leu Cys Glu Ile Lys Glu Lys Thr Val Leu Ser Ile Arg Gly Ala Gln
 85 90 95

Glu Glu Glu Pro Thr Asp Pro Gln Leu Met Arg Leu Asp Asn Met Leu
 100 105 110

Leu Ala Glu Gly Val Ala Gly Pro Glu Lys Gly Gly Gly Ser Ala Ala
 115 120 125

Ala Ala Ala Ala Ala Ala Ala Ser Gly Gly Ala Gly Ser Asp Asn Ser
 130 135 140

Val Glu His Ser Asp Tyr Arg Ala Lys Leu Ser Gln Ile Arg Gln Ile
 145 150 155 160

Tyr His Thr Glu Leu Glu Lys Tyr Glu Gln Ala Cys Asn Glu Phe Thr
 165 170 175

Thr His Val Met Asn Leu Leu Arg Glu Gln Ser Arg Thr Arg Pro Ile
 180 185 190

Ser Pro Lys Glu Ile Glu Arg Met Val Ser Ile Ile His Arg Lys Phe
 195 200 205

Ser Ser Ile Gln Met Gln Leu Lys Gln Ser Thr Cys Glu Ala Val Met
 210 215 220

Ile Leu Arg Ser Arg Phe Leu Asp Ala Arg Arg Lys Arg Arg Asn Phe
 225 230 235 240

Asn Lys Gln Ala Thr Glu Ile Leu Asn Glu Tyr Phe Tyr Ser His Leu
 245 250 255

Ser Asn Pro Tyr Pro Ser Glu Glu Ala Lys Glu Glu Leu Ala Lys Lys
 260 265 270

Cys Gly Ile Thr Val Ser Gln Val Ser Asn Trp Phe Gly Asn Lys Arg

275 280 285
 Ile Arg Tyr Lys Lys Asn Ile Gly Lys Phe Gln Glu Glu Ala Asn Ile
 290 295 300
 Tyr Ala Ala Lys Thr Ala Val Thr Ala Thr Asn Val Ser Ala His Gly
 305 310 315 320
 Ser Gln Ala Asn Ser Pro Ser Thr Pro Asn Ser Ala Gly Ser Ser Ser
 325 330 335
 Ser Phe Asn Met Ser Asn Ser Gly Asp Leu Phe Met Ser Val Gln Ser
 340 345 350
 Leu Asn Gly Asp Ser Tyr Gln Gly Ala Gln Val Gly Ala Asn Val Gln
 355 360 365
 Ser Gln Val Asp Thr Leu Arg His Val Ile Ser Gln Thr Gly Gly Tyr
 370 375 380
 Ser Asp Gly Leu Ala Ala Ser Gln Met Tyr Ser Pro Gln Gly Ile Ser
 385 390 395 400
 Ala Asn Gly Gly Trp Gln Asp Ala Thr Thr Pro Ser Ser Val Thr Ser
 405 410 415
 Pro Thr Glu Gly Pro Gly Ser Val His Ser Asp Thr Ser Asn
 420 425 430

 <210> 204
 <211> 384
 <212> PRT
 <213> Homo sapiens

 <400> 204
 Ala Arg Gly Val Ala Ser Met Thr Met Asn Val Ile Gln Thr Val Pro
 1 5 10 15
 Asn Leu Asp Trp Leu Ser Val Trp Ile Lys Ala Tyr Ala Phe Val His
 20 25 30
 Thr Gly Asp Asn Ser Arg Ala Ile Ser Thr Ile Cys Ser Leu Glu Lys
 35 40 45

Lys Ser Leu Leu Arg Asp Asn Val Asp Leu Leu Gly Ser Leu Ala Asp
50 55 60

Leu Tyr Phe Arg Ala Gly Asp Asn Lys Asn Ser Val Leu Lys Phe Glu
65 70 75 80

Gln Ala Gln Met Leu Asp Pro Tyr Leu Ile Lys Gly Met Asp Val Tyr
85 90 95

Gly Tyr Leu Leu Ala Arg Glu Gly Arg Leu Glu Asp Val Glu Asn Leu
100 105 110

Gly Cys Arg Leu Phe Asn Ile Ser Asp Gln His Ala Glu Pro Trp Val
115 120 125

Val Ser Gly Cys His Ser Phe Tyr Ser Lys Arg Tyr Ser Arg Ala Leu
130 135 140

Tyr Leu Gly Ala Lys Ala Ile Gln Leu Asn Ser Asn Ser Val Gln Ala
145 150 155 160

Leu Leu Leu Lys Gly Ala Ala Leu Arg Asn Met Gly Arg Val Gln Glu
165 170 175

Ala Ile Ile His Phe Arg Glu Ala Ile Arg Leu Ala Pro Cys Arg Leu
180 185 190

Asp Cys Tyr Glu Gly Leu Ile Glu Cys Tyr Leu Ala Ser Asn Ser Ile
195 200 205

Arg Glu Ala Met Val Met Ala Asn Asn Val Tyr Lys Thr Leu Gly Ala
210 215 220

Asn Ala Gln Thr Leu Thr Leu Leu Ala Thr Val Cys Leu Glu Asp Pro
225 230 235 240

Val Thr Gln Glu Lys Ala Lys Thr Leu Leu Asp Lys Ala Leu Thr Gln
245 250 255

Arg Pro Asp Tyr Ile Lys Ala Val Val Lys Lys Ala Glu Leu Leu Ser
260 265 270

Arg Glu Gln Lys Tyr Glu Asp Gly Ile Ala Leu Leu Arg Asn Ala Leu
 275 280 285

Ala Asn Gln Ser Asp Cys Val Leu His Arg Ile Leu Gly Asp Phe Leu
 290 295 300

Val Ala Val Asn Glu Tyr Gln Glu Ala Met Asp Gln Tyr Ser Ile Ala
 305 310 315 320

Leu Ser Leu Asp Pro Asn Asp Gln Lys Ser Leu Glu Gly Met Gln Lys
 325 330 335

Met Glu Lys Glu Glu Ser Pro Thr Asp Ala Thr Gln Glu Glu Asp Val
 340 345 350

Asp Asp Met Glu Gly Ser Gly Glu Glu Gly Asp Leu Glu Gly Ser Asp
 355 360 365

Ser Glu Ala Ala Gln Trp Ala Asp Gln Glu Gln Trp Phe Gly Met Gln
 370 375 380

<210> 205
 <211> 1659
 <212> PRT
 <213> Homo sapiens

<400> 205

Met Glu Ala Pro Ser Gly Ser Glu Pro Gly Gly Asp Gly Ala Gly Asp
 1 5 10 15

Cys Ala His Pro Asp Pro Arg Ala Pro Gly Ala Ala Ala Pro Ser Ser
 20 25 30

Gly Pro Gly Pro Cys Ala Ala Ala Arg Glu Ser Glu Arg Gln Leu Arg
 35 40 45

Leu Arg Leu Cys Val Leu Asn Glu Ile Leu Gly Thr Glu Arg Asp Tyr
 50 55 60

Val Gly Thr Leu Arg Phe Leu Gln Ser Ala Phe Leu His Arg Ile Arg
 65 70 75 80

Gln Asn Val Ala Asp Ser Val Glu Lys Gly Leu Thr Glu Glu Asn Val
 85 90 95

Lys Val Leu Phe Ser Asn Ile Glu Asp Ile Leu Glu Val His Lys Asp
 100 105 110

Phe Leu Ala Ala Leu Glu Tyr Cys Leu His Pro Glu Pro Gln Ser Gln
 115 120 125

His Glu Leu Gly Asn Val Phe Leu Lys Phe Lys Asp Lys Phe Cys Val
 130 135 140

Tyr Glu Glu Tyr Cys Ser Asn His Glu Lys Ala Leu Arg Leu Leu Val
 145 150 155 160

Glu Leu Asn Lys Ile Pro Thr Val Arg Ala Phe Leu Leu Ser Cys Met
 165 170 175

Leu Leu Gly Gly Arg Lys Thr Thr Asp Ile Pro Leu Glu Gly Tyr Leu
 180 185 190

Leu Ser Pro Ile Gln Arg Ile Cys Lys Tyr Pro Leu Leu Leu Lys Glu
 195 200 205

Leu Ala Lys Arg Thr Pro Gly Lys His Pro Asp His Pro Ala Val Gln
 210 215 220

Ser Ala Leu Gln Ala Met Lys Thr Val Cys Ser Asn Ile Asn Glu Thr
 225 230 235 240

Lys Arg Gln Met Glu Lys Leu Glu Ala Leu Glu Gln Leu Gln Ser His
 245 250 255

Ile Glu Gly Trp Glu Gly Ser Asn Leu Thr Asp Ile Cys Thr Gln Leu
 260 265 270

Leu Leu Gln Gly Thr Leu Leu Lys Ile Ser Ala Gly Asn Ile Gln Glu
 275 280 285

Arg Ala Phe Phe Leu Phe Asp Asn Leu Leu Val Tyr Cys Lys Arg Lys
 290 295 300

Ser Arg Val Thr Gly Ser Lys Lys Ser Thr Lys Arg Thr Lys Ser Ile
 305 310 315 320

Asn Gly Ser Leu Tyr Ile Phe Arg Gly Arg Ile Asn Thr Glu Val Met
 325 330 335

Glu Val Glu Asn Val Glu Asp Gly Thr Ala Asp Tyr His Ser Asn Gly
 340 345 350

Tyr Thr Val Thr Asn Gly Trp Lys Ile His Asn Thr Ala Lys Asn Lys
 355 360 365

Trp Phe Val Cys Met Ala Lys Thr Ala Glu Glu Lys Gln Lys Trp Leu
 370 375 380

Asp Ala Ile Ile Arg Glu Arg Glu Gln Arg Glu Ser Leu Lys Leu Gly
 385 390 395 400

Met Glu Arg Asp Ala Tyr Val Met Ile Ala Glu Lys Gly Glu Lys Leu
 405 410 415

Tyr His Met Met Met Asn Lys Lys Val Asn Leu Ile Lys Asp Arg Arg
 420 425 430

Arg Lys Leu Ser Thr Val Pro Lys Cys Phe Leu Gly Asn Glu Phe Val
 435 440 445

Ala Trp Leu Leu Glu Ile Gly Glu Ile Ser Lys Thr Glu Glu Gly Val
 450 455 460

Asn Leu Gly Gln Ala Leu Leu Glu Asn Gly Ile Ile His His Val Ser
 465 470 475 480

Asp Lys His Gln Phe Lys Asn Glu Gln Val Met Tyr Arg Phe Arg Tyr
 485 490 495

Asp Asp Gly Thr Tyr Lys Ala Arg Ser Glu Leu Glu Asp Ile Met Ser
 500 505 510

Lys Gly Val Arg Leu Tyr Cys Arg Leu His Ser Leu Tyr Thr Pro Val
 515 520 525

Ile Lys Asp Arg Asp Tyr His Leu Lys Thr Tyr Lys Ser Val Leu Pro
 530 535 540

Gly Ser Lys Leu Val Asp Trp Leu Leu Ala Gln Gly Asp Cys Gln Thr
 545 550 555 560

Arg Glu Glu Ala Val Ala Leu Gly Val Gly Leu Cys Asn Asn Gly Phe
 565 570 575

Met His His Val Leu Glu Lys Ser Glu Phe Arg Asp Glu Ser Gln Tyr
 580 585 590

Phe Arg Phe His Ala Asp Glu Glu Met Glu Gly Thr Ser Ser Lys Asn
 595 600 605

Lys Gln Leu Arg Asn Asp Phe Lys Leu Val Glu Asn Ile Leu Ala Lys
 610 615 620

Arg Leu Leu Ile Leu Pro Gln Glu Glu Asp Tyr Gly Phe Asp Ile Glu
 625 630 635 640

Glu Lys Asn Lys Ala Val Val Val Lys Ser Val Gln Arg Gly Ser Leu
 645 650 655

Ala Glu Val Ala Gly Leu Gln Val Gly Arg Lys Ile Tyr Ser Ile Asn
 660 665 670

Glu Asp Leu Val Phe Leu Arg Pro Phe Ser Glu Val Glu Ser Ile Leu
 675 680 685

Asn Gln Ser Phe Cys Ser Arg Arg Pro Leu Arg Leu Leu Val Ala Thr
 690 695 700

Lys Ala Lys Glu Ile Ile Lys Ile Pro Asp Gln Pro Asp Thr Leu Cys
 705 710 715 720

Phe Gln Ile Arg Gly Ala Ala Pro Pro Tyr Val Tyr Ala Val Gly Arg
 725 730 735

Gly Ser Glu Ala Met Ala Ala Gly Leu Cys Ala Gly Gln Cys Ile Leu
 740 745 750

Lys Val Asn Gly Ser Asn Val Met Asn Asp Gly Ala Pro Glu Val Leu
 755 760 765

Glu His Phe Gln Ala Phe Arg Ser Arg Arg Glu Glu Ala Leu Gly Leu

770 775 780
 Tyr Gln Trp Ile Tyr His Thr His Glu Asp Ala Gln Glu Ala Arg Ala
 785 790 795 800
 Ser Gln Glu Ala Ser Thr Glu Asp Pro Ser Gly Glu Gln Ala Gln Glu
 805 810 815
 Glu Asp Gln Ala Asp Ser Ala Phe Pro Leu Leu Ser Leu Gly Pro Arg
 820 825 830
 Leu Ser Leu Cys Glu Asp Ser Pro Met Val Thr Leu Thr Val Asp Asn
 835 840 845
 Val His Leu Glu His Gly Val Val Tyr Glu Tyr Val Ser Thr Ala Gly
 850 855 860
 Val Arg Cys His Val Leu Glu Lys Ile Val Glu Pro Arg Gly Cys Phe
 865 870 875 880
 Gly Leu Thr Ala Lys Ile Leu Glu Ala Phe Ala Ala Asn Asp Ser Val
 885 890 895
 Phe Val Glu Asn Cys Arg Arg Leu Met Ala Leu Ser Ser Ala Ile Val
 900 905 910
 Thr Met Pro His Phe Glu Phe Arg Asn Ile Cys Asp Thr Lys Leu Glu
 915 920 925
 Ser Ile Gly Gln Arg Ile Ala Cys Tyr Gln Glu Phe Ala Ala Gln Leu
 930 935 940
 Lys Ser Arg Val Ser Pro Pro Phe Lys Gln Ala Pro Leu Glu Pro His
 945 950 955 960
 Pro Leu Cys Gly Leu Asp Phe Cys Pro Thr Asn Cys His Ile Asn Leu
 965 970 975
 Met Glu Val Ser Tyr Pro Lys Thr Thr Pro Ser Val Gly Arg Ser Phe
 980 985 990
 Ser Ile Arg Phe Gly Arg Lys Pro Ser Leu Ile Gly Leu Asp Pro Glu
 995 1000 1005

Gln Gly	His Leu Asn Pro Met	Ser Tyr Thr Gln His	Cys Ile Thr
1010	1015	1020	
Thr Met	Ala Ala Pro Ser Trp	Lys Cys Leu Pro Ala	Ala Glu Gly
1025	1030	1035	
Asp Pro	Gln Gly Gln Gly Leu	His Asp Gly Ser Phe	Gly Pro Ala
1040	1045	1050	
Ser Gly	Thr Leu Gly Gln Glu	Asp Arg Gly Leu Ser	Phe Leu Leu
1055	1060	1065	
Lys Gln	Glu Asp Arg Glu Ile	Gln Asp Ala Tyr Leu	Gln Leu Phe
1070	1075	1080	
Thr Lys	Leu Asp Val Ala Leu	Lys Glu Met Lys Gln	Tyr Val Thr
1085	1090	1095	
Gln Ile	Asn Arg Leu Leu Ser	Thr Ile Thr Glu Pro	Thr Ser Gly
1100	1105	1110	
Gly Ser	Cys Asp Ala Ser Leu	Ala Glu Glu Ala Ser	Ser Leu Pro
1115	1120	1125	
Leu Val	Ser Glu Glu Ser Glu	Met Asp Arg Ser Asp	His Gly Gly
1130	1135	1140	
Ile Lys	Lys Val Cys Phe Lys	Val Ala Glu Glu Asp	Gln Glu Asp
1145	1150	1155	
Ser Gly	His Asp Thr Met Ser	Tyr Arg Asp Ser Tyr	Ser Glu Cys
1160	1165	1170	
Asn Ser	Asn Arg Asp Ser Val	Leu Ser Tyr Thr Ser	Val Arg Ser
1175	1180	1185	
Asn Ser	Ser Tyr Leu Gly Ser	Asp Glu Met Gly Ser	Gly Asp Glu
1190	1195	1200	
Leu Pro	Cys Asp Met Arg Ile	Pro Ser Asp Lys Gln	Asp Lys Leu
1205	1210	1215	

His Gly 1220	Cys Leu Glu	His Leu 1225	Phe Asn Gln Val	Asp 1230	Ser Ile Asn
Ala Leu 1235	Leu Lys Gly Pro	Val 1240	Met Ser Arg Ala	Phe 1245	Glu Glu Thr
Lys His 1250	Phe Pro Met Asn	His 1255	Ser Leu Gln Glu	Phe 1260	Lys Gln Lys
Glu Glu 1265	Cys Thr Ile Arg	Gly 1270	Arg Ser Leu Ile	Gln 1275	Ile Ser Ile
Gln Glu 1280	Asp Pro Trp Asn	Leu 1285	Pro Asn Ser Ile	Lys 1290	Thr Leu Val
Asp Asn 1295	Ile Gln Arg Tyr	Val 1300	Glu Asp Gly Lys	Asn 1305	Gln Leu Leu
Leu Ala 1310	Leu Leu Lys Cys	Thr 1315	Asp Thr Glu Leu	Gln 1320	Leu Arg Arg
Asp Ala 1325	Ile Phe Cys Gln	Ala 1330	Leu Val Ala Ala	Val 1335	Cys Thr Phe
Ser Glu 1340	Gln Leu Leu Ala	Ala 1345	Leu Gly Tyr Arg	Tyr 1350	Asn Asn Asn
Gly Glu 1355	Tyr Glu Glu Ser	Ser 1360	Arg Asp Ala Ser	Arg 1365	Lys Trp Leu
Glu Gln 1370	Val Ala Ala Thr	Gly 1375	Val Leu Leu His	Cys 1380	Gln Ser Leu
Leu Ser 1385	Pro Ala Thr Val	Lys 1390	Glu Glu Arg Thr	Met 1395	Leu Glu Asp
Ile Trp 1400	Val Thr Leu Ser	Glu 1405	Leu Asp Asn Val	Thr 1410	Phe Ser Phe
Lys Gln 1415	Leu Asp Glu Asn	Tyr 1420	Val Ala Asn Thr	Asn 1425	Val Phe Tyr

His Ile Glu Gly Ser Arg Gln Ala Leu Lys Val Ile Phe Tyr Leu
 1430 1435 1440

Asp Ser Tyr His Phe Ser Lys Leu Pro Ser Arg Leu Glu Gly Gly
 1445 1450 1455

Ala Ser Leu Arg Leu His Thr Ala Leu Phe Thr Lys Val Leu Glu
 1460 1465 1470

Asn Val Glu Gly Leu Pro Ser Pro Gly Ser Gln Ala Ala Glu Asp
 1475 1480 1485

Leu Gln Gln Asp Ile Asn Ala Gln Ser Leu Glu Lys Val Gln Gln
 1490 1495 1500

Tyr Tyr Arg Lys Leu Arg Ala Phe Tyr Leu Glu Arg Ser Asn Leu
 1505 1510 1515

Pro Thr Asp Ala Ser Thr Thr Ala Val Lys Ile Asp Gln Leu Ile
 1520 1525 1530

Arg Pro Ile Asn Ala Leu Asp Glu Leu Cys Arg Leu Met Lys Ser
 1535 1540 1545

Phe Val His Pro Lys Pro Gly Ala Ala Gly Ser Val Gly Ala Gly
 1550 1555 1560

Leu Ile Pro Ile Ser Ser Glu Leu Cys Tyr Arg Leu Gly Ala Cys
 1565 1570 1575

Gln Met Val Met Cys Gly Thr Gly Met Gln Arg Ser Thr Leu Ser
 1580 1585 1590

Val Ser Leu Glu Gln Ala Ala Ile Leu Ala Arg Ser His Gly Leu
 1595 1600 1605

Leu Pro Lys Cys Ile Met Gln Ala Thr Asp Ile Met Arg Lys Gln
 1610 1615 1620

Gly Pro Arg Val Glu Ile Leu Ala Lys Asn Leu Arg Val Lys Asp
 1625 1630 1635

Gln Met Pro Gln Gly Ala Pro Arg Leu Tyr Arg Leu Cys Gln Pro

1640

1645

1650

Pro Val Asp Gly Asp Leu
1655

<210> 206
<211> 175
<212> PRT
<213> Homo sapiens

<400> 206

Met Glu Lys Ile Pro Val Ser Ala Phe Leu Leu Leu Val Ala Leu Ser
1 5 10 15

Tyr Thr Leu Ala Arg Asp Thr Thr Val Lys Pro Gly Ala Lys Lys Asp
20 25 30

Thr Lys Asp Ser Arg Pro Lys Leu Pro Gln Thr Leu Ser Arg Gly Trp
35 40 45

Gly Asp Gln Leu Ile Trp Thr Gln Thr Tyr Glu Glu Ala Leu Tyr Lys
50 55 60

Ser Lys Thr Ser Asn Lys Pro Leu Met Ile Ile His His Leu Asp Glu
65 70 75 80

Cys Pro His Ser Gln Ala Leu Lys Lys Val Phe Ala Glu Asn Lys Glu
85 90 95

Ile Gln Lys Leu Ala Glu Gln Phe Val Leu Leu Asn Leu Val Tyr Glu
100 105 110

Thr Thr Asp Lys His Leu Ser Pro Asp Gly Gln Tyr Val Pro Arg Ile
115 120 125

Met Phe Val Asp Pro Ser Leu Thr Val Arg Ala Asp Ile Thr Gly Arg
130 135 140

Tyr Ser Asn Arg Leu Tyr Ala Tyr Glu Pro Ala Asp Thr Ala Leu Leu
145 150 155 160

Leu Asp Asn Met Lys Lys Ala Leu Lys Leu Leu Lys Thr Glu Leu
165 170 175

<210> 207
 <211> 196
 <212> PRT
 <213> Homo sapiens

<400> 207

Met Ala Ala Ile Arg Lys Lys Leu Val Val Val Gly Asp Gly Ala Cys
 1 5 10 15

Gly Lys Thr Cys Leu Leu Ile Val Phe Ser Lys Asp Glu Phe Pro Glu
 20 25 30

Val Tyr Val Pro Thr Val Phe Glu Asn Tyr Val Ala Asp Ile Glu Val
 35 40 45

Asp Gly Lys Gln Val Glu Leu Ala Leu Trp Asp Thr Ala Gly Gln Glu
 50 55 60

Asp Tyr Asp Arg Leu Arg Pro Leu Ser Tyr Pro Asp Thr Asp Val Ile
 65 70 75 80

Leu Met Cys Phe Ser Val Asp Ser Pro Asp Ser Leu Glu Asn Ile Pro
 85 90 95

Glu Lys Trp Val Pro Glu Val Lys His Phe Cys Pro Asn Val Pro Ile
 100 105 110

Ile Leu Val Ala Asn Lys Lys Asp Leu Arg Ser Asp Glu His Val Arg
 115 120 125

Thr Glu Leu Ala Arg Met Lys Gln Glu Pro Val Arg Thr Asp Asp Gly
 130 135 140

Arg Ala Met Ala Val Arg Ile Gln Ala Tyr Asp Tyr Leu Glu Cys Ser
 145 150 155 160

Ala Lys Thr Lys Glu Gly Val Arg Glu Val Phe Glu Thr Ala Thr Arg
 165 170 175

Ala Ala Leu Gln Lys Arg Tyr Gly Ser Gln Asn Gly Cys Ile Asn Cys
 180 185 190

Cys Lys Val Leu

195

<210> 208
 <211> 291
 <212> PRT
 <213> Homo sapiens

<400> 208

Met Glu Lys Leu Ala Ala Ser Thr Glu Pro Gln Gly Pro Arg Pro Val
 1 5 10 15

Leu Gly Arg Glu Ser Val Gln Val Pro Asp Asp Gln Asp Phe Arg Ser
 20 25 30

Phe Arg Ser Glu Cys Glu Ala Glu Val Gly Trp Asn Leu Thr Tyr Ser
 35 40 45

Arg Ala Gly Val Ser Val Trp Val Gln Ala Val Glu Met Asp Arg Thr
 50 55 60

Leu His Lys Ile Lys Cys Arg Met Glu Cys Cys Asp Val Pro Ala Glu
 65 70 75 80

Thr Leu Tyr Asp Val Leu His Asp Ile Glu Tyr Arg Lys Lys Trp Asp
 85 90 95

Ser Asn Val Ile Glu Thr Phe Asp Ile Ala Arg Leu Thr Val Asn Ala
 100 105 110

Asp Val Gly Tyr Tyr Ser Trp Arg Cys Pro Lys Pro Leu Lys Asn Arg
 115 120 125

Asp Val Ile Thr Leu Arg Ser Trp Leu Pro Met Gly Ala Asp Tyr Ile
 130 135 140

Ile Met Asn Tyr Ser Val Lys His Pro Lys Tyr Pro Pro Arg Lys Asp
 145 150 155 160

Leu Val Arg Ala Val Ser Ile Gln Thr Gly Tyr Leu Ile Gln Ser Thr
 165 170 175

Gly Pro Lys Ser Cys Val Ile Thr Tyr Leu Ala Gln Val Asp Pro Lys
 180 185 190

Gly Ser Leu Pro Lys Trp Val Val Asn Lys Ser Ser Gln Phe Leu Ala
 195 200 205

Pro Lys Ala Met Lys Lys Met Tyr Lys Ala Cys Leu Lys Tyr Pro Glu
 210 215 220

Trp Lys Gln Lys His Leu Pro His Phe Lys Pro Trp Leu His Pro Glu
 225 230 235 240

Gln Ser Pro Leu Pro Ser Leu Ala Leu Ser Glu Leu Ser Val Gln His
 245 250 255

Ala Asp Ser Leu Glu Asn Ile Asp Glu Ser Ala Val Ala Glu Ser Arg
 260 265 270

Glu Glu Arg Met Gly Gly Ala Gly Gly Glu Gly Ser Asp Asp Asp Thr
 275 280 285

Ser Leu Thr
 290

<210> 209
 <211> 358
 <212> PRT
 <213> Homo sapiens

<400> 209

Met Ser Ala Asp Ala Ala Ala Gly Ala Pro Leu Pro Arg Leu Cys Cys
 1 5 10 15

Leu Glu Lys Gly Pro Asn Gly Tyr Gly Phe His Leu His Gly Glu Lys
 20 25 30

Gly Lys Leu Gly Gln Tyr Ile Arg Leu Val Glu Pro Gly Ser Pro Ala
 35 40 45

Glu Lys Ala Gly Leu Leu Ala Gly Asp Arg Leu Val Glu Val Asn Gly
 50 55 60

Glu Asn Val Glu Lys Glu Thr His Gln Gln Val Val Ser Arg Ile Arg
 65 70 75 80

Ala Ala Leu Asn Ala Val Arg Leu Leu Val Val Asp Pro Glu Thr Asp

356

Pro Ala Ser Ser Val Ser Ser Ser Pro Ser Pro Pro Phe Gly His Ser
145 150 155 160

Ala Ala Val Ser Pro Thr Phe Met Pro Arg Ser Thr Gln Pro Leu Thr
165 170 175

Phe Thr Thr Ala Thr Phe Ala Ala Thr Lys Phe Gly Ser Thr Lys Met
180 185 190

Lys Asn Ser Gly Arg Ser Asn Lys Val Ala Arg Thr Ser Pro Ile Asn
195 200 205

Leu Gly Leu Asn Val Asn Asp Leu Leu Lys Gln Lys Ala Ile Ser Ser
210 215 220

Ser Met His Ser Leu Tyr Gly Leu Gly Leu Gly Ser Gln Gln Gln Pro
225 230 235 240

Gln Gln Gln Gln Gln Pro Ala Gln Pro Pro Pro Pro Pro Pro Pro
245 250 255

Gln Gln Gln Gln Gln Gln Lys Thr Ser Ala Leu Ser Pro Asn Ala Lys
260 265 270

Glu Phe Ile Phe Pro Asn Met Gln Gly Gln Gly Ser Ser Thr Asn Gly
275 280 285

Met Phe Pro Gly Asp Ser Pro Leu Asn Leu Ser Pro Leu Gln Tyr Ser
290 295 300

Asn Ala Phe Asp Val Phe Ala Ala Tyr Gly Gly Leu Asn Glu Lys Ser
305 310 315 320

Phe Val Asp Gly Leu Asn Phe Ser Leu Asn Asn Met Gln Tyr Ser Asn
325 330 335

Gln Gln Phe Gln Pro Val Met Ala Asn
340 345

<210> 211
<211> 84
<212> PRT
<213> Homo sapiens

<400> 211

Met Ala Thr Met Glu Asn Lys Val Ile Cys Ala Leu Val Leu Val Ser
 1 5 10 15

Met Leu Ala Leu Gly Thr Leu Ala Glu Ala Gln Thr Glu Thr Cys Thr
 20 25 30

Val Ala Pro Arg Glu Arg Gln Asn Cys Gly Phe Pro Gly Val Thr Pro
 35 40 45

Ser Gln Cys Ala Asn Lys Gly Cys Cys Phe Asp Asp Thr Val Arg Gly
 50 55 60

Val Pro Trp Cys Phe Tyr Pro Asn Thr Ile Asp Val Pro Pro Glu Glu
 65 70 75 80

Glu Cys Glu Phe

<210> 212

<211> 522

<212> PRT

<213> Homo sapiens

<400> 212

Gly Phe Leu Pro Ala Thr Lys Asn Leu Leu Asn Glu Lys Asn His Gly
 1 5 10 15

Val Leu His Thr Ser Val Val Leu Leu Thr Glu Met Cys Glu Arg Ser
 20 25 30

Pro Asp Met Leu Ala His Phe Arg Glu Asn Glu Lys Leu Val Pro Gln
 35 40 45

Leu Val Arg Ile Leu Lys Asn Leu Ile Met Ser Gly Tyr Ser Pro Gly
 50 55 60

His Asp Val Ser Gly Ile Ser Asp Pro Phe Leu Gln Val Arg Ile Leu
 65 70 75 80

Arg Leu Leu Arg Ile Leu Gly Arg Asn Asp Asp Asp Ser Ser Glu Ala
 85 90 95

Met Asn Asp Ile Leu Ala Gln Val Ala Thr Asn Thr Glu Thr Ser Lys
 100 105 110

Asn Val Gly Asn Ala Ile Leu Tyr Glu Thr Val Leu Thr Ile Met Asp
 115 120 125

Ile Lys Ser Glu Ser Gly Leu Arg Val Leu Ala Ile Asn Ile Leu Gly
 130 135 140

Arg Phe Leu Leu Asn Asn Asp Lys Asn Ile Arg Tyr Val Ala Leu Thr
 145 150 155 160

Ser Leu Leu Lys Thr Val Gln Thr Asp His Asn Ala Val Gln Arg His
 165 170 175

Arg Ser Thr Ile Val Asp Cys Leu Lys Asp Leu Asp Val Ser Ile Lys
 180 185 190

Arg Arg Ala Met Glu Leu Ser Phe Ala Leu Val Asn Gly Asn Asn Ile
 195 200 205

Arg Gly Met Met Lys Glu Leu Leu Tyr Phe Leu Asp Ser Cys Glu Pro
 210 215 220

Glu Phe Lys Ala Asp Cys Ala Ser Gly Ile Phe Leu Ala Ala Glu Lys
 225 230 235 240

Tyr Ala Pro Ser Lys Arg Trp His Ile Asp Thr Ile Met Arg Val Leu
 245 250 255

Thr Thr Ala Gly Ser Tyr Val Arg Asp Asp Ala Val Pro Asn Leu Ile
 260 265 270

Gln Leu Ile Thr Asn Ser Val Glu Met His Ala Tyr Thr Val Gln Arg
 275 280 285

Leu Tyr Lys Ala Ile Leu Gly Asp Tyr Ser Gln Gln Pro Leu Val Gln
 290 295 300

Val Ala Ala Trp Cys Ile Gly Glu Tyr Gly Asp Leu Leu Val Ser Gly
 305 310 315 320

Gln Cys Glu Glu Glu Glu Pro Ile Gln Val Thr Glu Asp Glu Val Leu


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<210> 213
<211> 1704
<212> PRT
<213> Homo sapiens

<400> 213
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Met Ala Val Leu Arg Gln Leu Ala Leu Leu Leu Trp Lys Asn Tyr Thr
1 5 10 15

Leu Gln Lys Arg Lys Val Leu Val Thr Val Leu Glu Leu Phe Leu Pro
20 25 30

Leu Leu Phe Pro Gly Ile Leu Ile Trp Leu Arg Leu Lys Ile Gln Ser
35 40 45

Glu Asn Val Pro Asn Ala Thr Ile Tyr Pro Gly Gln Ser Ile Gln Glu
50 55 60

Leu Pro Leu Phe Phe Thr Phe Pro Pro Pro Gly Asp Thr Trp Glu Leu
65 70 75 80

Ala Tyr Ile Pro Ser His Ser Asp Ala Ala Lys Thr Val Thr Glu Thr
85 90 95

Val Arg Arg Ala Leu Val Ile Asn Met Arg Val Arg Gly Phe Pro Ser
100 105 110

Glu Lys Asp Phe Glu Asp Tyr Ile Arg Tyr Asp Asn Cys Ser Ser Ser
115 120 125

Val Leu Ala Ala Val Val Phe Glu His Pro Phe Asn His Ser Lys Glu
130 135 140

Pro Leu Pro Leu Ala Val Lys Tyr His Leu Arg Phe Ser Tyr Thr Arg
145 150 155 160

Arg Asn Tyr Met Trp Thr Gln Thr Gly Ser Phe Phe Leu Lys Glu Thr
165 170 175

Glu Gly Trp His Thr Thr Ser Leu Phe Pro Leu Phe Pro Asn Pro Gly
180 185 190

Pro Arg Glu Leu Thr Ser Pro Asp Gly Gly Glu Pro Gly Tyr Ile Arg
195 200 205

Glu Gly Phe Leu Ala Val Gln His Ala Val Asp Arg Ala Ile Met Glu
210 215 220

Tyr His Ala Asp Ala Ala Thr Arg Gln Leu Phe Gln Arg Leu Thr Val
225 230 235 240

Thr Ile Lys Arg Phe Pro Tyr Pro Pro Phe Ile Ala Asp Pro Phe Leu
245 250 255

Val Ala Ile Gln Tyr Gln Leu Pro Leu Leu Leu Leu Leu Ser Phe Thr
260 265 270

Tyr Thr Ala Leu Thr Ile Ala Arg Ala Val Val Gln Glu Lys Glu Arg
275 280 285

Arg Leu Lys Glu Tyr Met Arg Met Met Gly Leu Ser Ser Trp Leu His
290 295 300

Trp Ser Ala Trp Phe Leu Leu Phe Phe Leu Phe Leu Leu Ile Ala Ala
305 310 315 320

Ser Phe Met Thr Leu Leu Phe Cys Val Lys Val Lys Pro Asn Val Ala
325 330 335

Val Leu Ser Arg Ser Asp Pro Ser Leu Val Leu Ala Phe Leu Leu Cys
340 345 350

Phe Ala Ile Ser Thr Ile Ser Phe Ser Phe Met Val Ser Thr Phe Phe
355 360 365

Ser Lys Ala Asn Met Ala Ala Ala Phe Gly Gly Phe Leu Tyr Phe Phe
370 375 380

Thr Tyr Ile Pro Tyr Phe Phe Val Ala Pro Arg Tyr Asn Trp Met Thr
385 390 395 400

Leu Ser Gln Lys Leu Cys Ser Cys Leu Leu Ser Asn Val Ala Met Ala
405 410 415

Met Gly Ala Gln Leu Ile Gly Lys Phe Glu Ala Lys Gly Met Gly Ile
420 425 430

Gln Trp Arg Asp Leu Leu Ser Pro Val Asn Val Asp Asp Asp Phe Cys
435 440 445

Phe Gly Gln Val Leu Gly Met Leu Leu Leu Asp Ser Val Leu Tyr Gly

450 455 460
 Leu Val Thr Trp Tyr Met Glu Ala Val Phe Pro Gly Gln Phe Gly Val
 465 470 475 480
 Pro Gln Pro Trp Tyr Phe Phe Ile Met Pro Ser Tyr Trp Cys Gly Lys
 485 490 495
 Pro Arg Ala Val Ala Gly Lys Glu Glu Glu Asp Ser Asp Pro Glu Lys
 500 505 510
 Ala Leu Arg Asn Glu Tyr Phe Glu Ala Glu Pro Glu Asp Leu Val Ala
 515 520 525
 Gly Ile Lys Ile Lys His Leu Ser Lys Val Phe Arg Val Gly Asn Lys
 530 535 540
 Asp Arg Ala Ala Val Arg Asp Leu Asn Leu Asn Leu Tyr Glu Gly Gln
 545 550 555 560
 Ile Thr Val Leu Leu Gly His Asn Gly Ala Gly Lys Thr Thr Thr Leu
 565 570 575
 Ser Met Leu Thr Gly Leu Phe Pro Pro Thr Ser Gly Arg Ala Tyr Ile
 580 585 590
 Ser Gly Tyr Glu Ile Ser Gln Asp Met Val Gln Ile Arg Lys Ser Leu
 595 600 605
 Gly Leu Cys Pro Gln His Asp Ile Leu Phe Asp Asn Leu Thr Val Ala
 610 615 620
 Glu His Leu Tyr Phe Tyr Ala Gln Leu Lys Gly Leu Ser Arg Gln Lys
 625 630 635 640
 Cys Pro Glu Glu Val Lys Gln Met Leu His Ile Ile Gly Leu Glu Asp
 645 650 655
 Lys Trp Asn Ser Arg Ser Arg Phe Leu Ser Gly Gly Met Arg Arg Lys
 660 665 670
 Leu Ser Ile Gly Ile Ala Leu Ile Ala Gly Ser Lys Val Leu Ile Leu
 675 680 685

Asp Glu Pro Thr Ser Gly Met Asp Ala Ile Ser Arg Arg Ala Ile Trp
 690 695 700

Asp Leu Leu Gln Arg Gln Lys Ser Asp Arg Thr Ile Val Leu Thr Thr
 705 710 715 720

His Phe Met Asp Glu Ala Asp Leu Leu Gly Asp Arg Ile Ala Ile Met
 725 730 735

Ala Lys Gly Glu Leu Gln Cys Cys Gly Ser Ser Leu Phe Leu Lys Gln
 740 745 750

Lys Tyr Gly Ala Gly Tyr His Met Thr Leu Val Lys Glu Pro His Cys
 755 760 765

Asn Pro Glu Asp Ile Ser Gln Leu Val His His His Val Pro Asn Ala
 770 775 780

Thr Leu Glu Ser Ser Ala Gly Ala Glu Leu Ser Phe Ile Leu Pro Arg
 785 790 795 800

Glu Ser Thr His Arg Phe Glu Gly Leu Phe Ala Lys Leu Glu Lys Lys
 805 810 815

Gln Lys Glu Leu Gly Ile Ala Ser Phe Gly Ala Ser Ile Thr Thr Met
 820 825 830

Glu Glu Val Phe Leu Arg Val Gly Lys Leu Val Asp Ser Ser Met Asp
 835 840 845

Ile Gln Ala Ile Gln Leu Pro Ala Leu Gln Tyr Gln His Glu Arg Arg
 850 855 860

Ala Ser Asp Trp Ala Val Asp Ser Asn Leu Cys Gly Ala Met Asp Pro
 865 870 875 880

Ser Asp Gly Ile Gly Ala Leu Ile Glu Glu Glu Arg Thr Ala Val Lys
 885 890 895

Leu Asn Thr Gly Leu Ala Leu His Cys Gln Gln Phe Trp Ala Met Phe
 900 905 910

Leu Lys Lys Ala Ala Tyr Ser Trp Arg Glu Trp Lys Met Val Ala Ala
 915 920 925

Gln Val Leu Val Pro Leu Thr Cys Val Thr Leu Ala Leu Leu Ala Ile
 930 935 940

Asn Tyr Ser Ser Glu Leu Phe Asp Asp Pro Met Leu Arg Leu Thr Leu
 945 950 955 960

Gly Glu Tyr Gly Arg Thr Val Val Pro Phe Ser Val Pro Gly Thr Ser
 965 970 975

Gln Leu Gly Gln Gln Leu Ser Glu His Leu Lys Asp Ala Leu Gln Ala
 980 985 990

Glu Gly Gln Glu Pro Arg Glu Val Leu Gly Asp Leu Glu Glu Phe Leu
 995 1000 1005

Ile Phe Arg Ala Ser Val Glu Gly Gly Gly Phe Asn Glu Arg Cys
 1010 1015 1020

Leu Val Ala Ala Ser Phe Arg Asp Val Gly Glu Arg Thr Val Val
 1025 1030 1035

Asn Ala Leu Phe Asn Asn Gln Ala Tyr His Ser Pro Ala Thr Ala
 1040 1045 1050

Leu Ala Val Val Asp Asn Leu Leu Phe Lys Leu Leu Cys Gly Pro
 1055 1060 1065

His Ala Ser Ile Val Val Ser Asn Phe Pro Gln Pro Arg Ser Ala
 1070 1075 1080

Leu Gln Ala Ala Lys Asp Gln Phe Asn Glu Gly Arg Lys Gly Phe
 1085 1090 1095

Asp Ile Ala Leu Asn Leu Leu Phe Ala Met Ala Phe Leu Ala Ser
 1100 1105 1110

Thr Phe Ser Ile Leu Ala Val Ser Glu Arg Ala Val Gln Ala Lys
 1115 1120 1125

His Val	Gln Phe Val Ser Gly	Val His Val Ala Ser	Phe Trp Leu
1130	1135	1140	
Ser Ala	Leu Leu Trp Asp Leu	Ile Ser Phe Leu Ile	Pro Ser Leu
1145	1150	1155	
Leu Leu	Leu Val Val Phe Lys	Ala Phe Asp Val Arg	Ala Phe Thr
1160	1165	1170	
Arg Asp	Gly His Met Ala Asp	Thr Leu Leu Leu Leu	Leu Leu Tyr
1175	1180	1185	
Gly Trp	Ala Ile Ile Pro Leu	Met Tyr Leu Met Asn	Phe Phe Phe
1190	1195	1200	
Leu Gly	Ala Ala Thr Ala Tyr	Thr Arg Leu Thr Ile	Phe Asn Ile
1205	1210	1215	
Leu Ser	Gly Ile Ala Thr Phe	Leu Met Val Thr Ile	Met Arg Ile
1220	1225	1230	
Pro Ala	Val Lys Leu Glu Glu	Leu Ser Lys Thr Leu	Asp His Val
1235	1240	1245	
Phe Leu	Val Leu Pro Asn His	Cys Leu Gly Met Ala	Val Ser Ser
1250	1255	1260	
Phe Tyr	Glu Asn Tyr Glu Thr	Arg Arg Tyr Cys Thr	Ser Ser Glu
1265	1270	1275	
Val Ala	Ala His Tyr Cys Lys	Lys Tyr Asn Ile Gln	Tyr Gln Glu
1280	1285	1290	
Asn Phe	Tyr Ala Trp Ser Ala	Pro Gly Val Gly Arg	Phe Val Ala
1295	1300	1305	
Ser Met	Ala Ala Ser Gly Cys	Ala Tyr Leu Ile Leu	Leu Phe Leu
1310	1315	1320	
Ile Glu	Thr Asn Leu Leu Gln	Arg Leu Arg Gly Ile	Leu Cys Ala
1325	1330	1335	
Leu Arg	Arg Arg Arg Thr Leu	Thr Glu Leu Tyr Thr	Arg Met Pro

1340	1345	1350
Val Leu Pro Glu Asp Gln Asp	Val Ala Asp Glu Arg	Thr Arg Ile
1355	1360	1365
Leu Ala Pro Ser Pro Asp Ser	Leu Leu His Thr Pro	Leu Ile Ile
1370	1375	1380
Lys Glu Leu Ser Lys Val Tyr	Glu Gln Arg Val Pro	Leu Leu Ala
1385	1390	1395
Val Asp Arg Leu Ser Leu Ala	Val Gln Lys Gly Glu	Cys Phe Gly
1400	1405	1410
Leu Leu Gly Phe Asn Gly Ala	Gly Lys Thr Thr Thr	Phe Lys Met
1415	1420	1425
Leu Thr Gly Glu Glu Ser Leu	Thr Ser Gly Asp Ala	Phe Val Gly
1430	1435	1440
Gly His Arg Ile Ser Ser Asp	Val Gly Lys Val Arg	Gln Arg Ile
1445	1450	1455
Gly Tyr Cys Pro Gln Phe Asp	Ala Leu Leu Asp His	Met Thr Gly
1460	1465	1470
Arg Glu Met Leu Val Met Tyr	Ala Arg Leu Arg Gly	Ile Pro Glu
1475	1480	1485
Arg His Ile Gly Ala Cys Val	Glu Asn Thr Leu Arg	Gly Leu Leu
1490	1495	1500
Leu Glu Pro His Ala Asn Lys	Leu Val Arg Thr Tyr	Ser Gly Gly
1505	1510	1515
Asn Lys Arg Lys Leu Ser Thr	Gly Ile Ala Leu Ile	Gly Glu Pro
1520	1525	1530
Ala Val Ile Phe Leu Asp Glu	Pro Ser Thr Gly Met	Asp Pro Val
1535	1540	1545
Ala Arg Arg Leu Leu Trp Asp	Thr Val Ala Arg Ala	Arg Glu Ser
1550	1555	1560

Gly Lys Ala Ile Ile Ile Thr Ser His Ser Met Glu Glu Cys Glu
1565 1570 1575

Ala Leu Cys Thr Arg Leu Ala Ile Met Val Gln Gly Gln Phe Lys
1580 1585 1590

Cys Leu Gly Ser Pro Gln His Leu Lys Ser Lys Phe Gly Ser Gly
1595 1600 1605

Tyr Ser Leu Arg Ala Lys Val Gln Ser Glu Gly Gln Gln Glu Ala
1610 1615 1620

Leu Glu Glu Phe Lys Ala Phe Val Asp Leu Thr Phe Pro Gly Ser
1625 1630 1635

Val Leu Glu Asp Glu His Gln Gly Met Val His Tyr His Leu Pro
1640 1645 1650

Gly Arg Asp Leu Ser Trp Ala Lys Val Phe Gly Ile Leu Glu Lys
1655 1660 1665

Ala Lys Glu Lys Tyr Gly Val Asp Asp Tyr Ser Val Ser Gln Ile
1670 1675 1680

Ser Leu Glu Gln Val Phe Leu Ser Phe Ala His Leu Gln Pro Pro
1685 1690 1695

Thr Ala Glu Glu Gly Arg
1700

<210> 214
<211> 674
<212> PRT
<213> Homo sapiens

<400> 214

Met Ala Ala Phe Ser Val Gly Thr Ala Met Asn Ala Ser Ser Tyr Ser
1 5 10 15

Ala Glu Met Thr Glu Pro Lys Ser Val Cys Val Ser Val Asp Glu Val
20 25 30

Val Ser Ser Asn Met Glu Ala Thr Glu Thr Asp Leu Leu Asn Gly His
 35 40 45

Leu Lys Lys Val Asp Asn Asn Leu Thr Glu Ala Gln Arg Phe Ser Ser
 50 55 60

Leu Pro Arg Arg Ala Ala Val Asn Ile Glu Phe Arg Asp Leu Ser Tyr
 65 70 75 80

Ser Val Pro Glu Gly Pro Trp Trp Arg Lys Lys Gly Tyr Lys Thr Leu
 85 90 95

Leu Lys Gly Ile Ser Gly Lys Phe Asn Ser Gly Glu Leu Val Ala Ile
 100 105 110

Met Gly Pro Ser Gly Ala Gly Lys Ser Thr Leu Met Asn Ile Leu Ala
 115 120 125

Gly Tyr Arg Glu Thr Gly Met Lys Gly Ala Val Leu Ile Asn Gly Leu
 130 135 140

Pro Arg Asp Leu Arg Cys Phe Arg Lys Val Ser Cys Tyr Ile Met Gln
 145 150 155 160

Asp Asp Met Leu Leu Pro His Leu Thr Val Gln Glu Ala Met Met Val
 165 170 175

Ser Ala His Leu Lys Leu Gln Glu Lys Asp Glu Gly Arg Arg Glu Met
 180 185 190

Val Lys Glu Ile Leu Thr Ala Leu Gly Leu Leu Ser Cys Ala Asn Thr
 195 200 205

Arg Thr Gly Ser Leu Ser Gly Gly Gln Arg Lys Arg Leu Ala Ile Ala
 210 215 220

Leu Glu Leu Val Asn Asn Pro Pro Val Met Phe Phe Asp Glu Pro Thr
 225 230 235 240

Ser Gly Leu Asp Ser Ala Ser Cys Phe Gln Val Val Ser Leu Met Lys
 245 250 255

Gly Leu Ala Gln Gly Gly Arg Ser Ile Ile Cys Thr Ile His Gln Pro

260	265	270
Ser Ala Lys Leu Phe Glu Leu Phe Asp Gln Leu Tyr Val Leu Ser Gln		
275	280	285
Gly Gln Cys Val Tyr Arg Gly Lys Val Cys Asn Leu Val Pro Tyr Leu		
290	295	300
Arg Asp Leu Gly Leu Asn Cys Pro Thr Tyr His Asn Pro Ala Asp Phe		
305	310	315 320
Val Met Glu Val Ala Ser Gly Glu Tyr Gly Asp Gln Asn Ser Arg Leu		
325	330	335
Val Arg Ala Val Arg Glu Gly Met Cys Asp Ser Asp His Lys Arg Asp		
340	345	350
Leu Gly Gly Asp Ala Glu Val Asn Pro Phe Leu Trp His Arg Pro Ser		
355	360	365
Glu Glu Val Lys Gln Thr Lys Arg Leu Lys Gly Leu Arg Lys Asp Ser		
370	375	380
Ser Ser Met Glu Gly Cys His Ser Phe Ser Ala Ser Cys Leu Thr Gln		
385	390	395 400
Phe Cys Ile Leu Phe Lys Arg Thr Phe Leu Ser Ile Met Arg Asp Ser		
405	410	415
Val Leu Thr His Leu Arg Ile Thr Ser His Ile Gly Ile Gly Leu Leu		
420	425	430
Ile Gly Leu Leu Tyr Leu Gly Ile Gly Asn Glu Thr Lys Lys Val Leu		
435	440	445
Ser Asn Ser Gly Phe Leu Phe Phe Ser Met Leu Phe Leu Met Phe Ala		
450	455	460
Ala Leu Met Pro Thr Val Leu Thr Phe Pro Leu Glu Met Gly Val Phe		
465	470	475 480
Leu Arg Glu His Leu Asn Tyr Trp Tyr Ser Leu Lys Ala Tyr Tyr Leu		
485	490	495

Ala Lys Thr Met Ala Asp Val Pro Phe Gln Ile Met Phe Pro Val Ala
 500 505 510

Tyr Cys Ser Ile Val Tyr Trp Met Thr Ser Gln Pro Ser Asp Ala Val
 515 520 525

Arg Phe Val Leu Phe Ala Ala Leu Gly Thr Met Thr Ser Leu Val Ala
 530 535 540

Gln Ser Leu Gly Leu Leu Ile Gly Ala Ala Ser Thr Ser Leu Gln Val
 545 550 555 560

Ala Thr Phe Val Gly Pro Val Thr Ala Ile Pro Val Leu Leu Phe Ser
 565 570 575

Gly Phe Phe Val Ser Phe Asp Thr Ile Pro Thr Tyr Leu Gln Trp Met
 580 585 590

Ser Tyr Ile Ser Tyr Val Arg Tyr Gly Phe Glu Gly Val Ile Leu Ser
 595 600 605

Ile Tyr Gly Leu Asp Arg Glu Asp Leu His Cys Asp Ile Asp Glu Thr
 610 615 620

Cys His Phe Gln Lys Ser Glu Ala Ile Leu Arg Glu Leu Asp Val Glu
 625 630 635 640

Asn Ala Lys Leu Tyr Leu Asp Phe Ile Val Leu Gly Ile Phe Phe Ile
 645 650 655

Ser Leu Arg Leu Ile Ala Tyr Leu Val Leu Arg Tyr Lys Ile Arg Ala
 660 665 670

Glu Arg

<210> 215
 <211> 149
 <212> PRT
 <213> Homo sapiens
 <400> 215

Met Ala Asp Gln Leu Thr Glu Glu Gln Ile Ala Glu Phe Lys Glu Ala
1 5 10 15

Phe Ser Leu Phe Asp Lys Asp Gly Asp Gly Thr Ile Thr Thr Lys Glu
20 25 30

Leu Gly Thr Val Met Arg Ser Leu Gly Gln Asn Pro Thr Glu Ala Glu
35 40 45

Leu Gln Asp Met Ile Asn Glu Val Asp Ala Asp Gly Asn Gly Thr Ile
50 55 60

Asp Phe Pro Glu Phe Leu Thr Met Met Ala Arg Lys Met Lys Asp Thr
65 70 75 80

Asp Ser Glu Glu Glu Ile Arg Glu Ala Phe Arg Val Phe Asp Lys Asp
85 90 95

Gly Asn Gly Tyr Ile Ser Ala Ala Glu Leu Arg His Val Met Thr Asn
100 105 110

Leu Gly Glu Lys Leu Thr Asp Glu Glu Val Asp Glu Met Ile Arg Glu
115 120 125

Ala Asp Ile Asp Gly Asp Gly Gln Val Asn Tyr Glu Glu Phe Val Gln
130 135 140

Met Met Thr Ala Lys
145

<210> 216
<211> 354
<212> PRT
<213> Homo sapiens

<400> 216

Met Pro Arg Arg Ser Leu His Ala Ala Ala Val Leu Leu Leu Val Ile
1 5 10 15

Leu Lys Glu Gln Pro Ser Ser Pro Ala Pro Val Asn Gly Ser Lys Trp
20 25 30

Thr Tyr Phe Gly Pro Asp Gly Glu Asn Ser Trp Ser Lys Lys Tyr Pro
35 40 45

Ser Cys Gly Gly Leu Leu Gln Ser Pro Ile Asp Leu His Ser Asp Ile
50 55 60

Leu Gln Tyr Asp Ala Ser Leu Thr Pro Leu Glu Phe Gln Gly Tyr Asn
65 70 75 80

Leu Ser Ala Asn Lys Gln Phe Leu Leu Thr Asn Asn Gly His Ser Val
85 90 95

Lys Leu Asn Leu Pro Ser Asp Met His Ile Gln Gly Leu Gln Ser Arg
100 105 110

Tyr Ser Ala Thr Gln Leu His Leu His Trp Gly Asn Pro Asn Asp Pro
115 120 125

His Gly Ser Glu His Thr Val Ser Gly Gln His Phe Ala Ala Glu Leu
130 135 140

His Ile Val His Tyr Asn Ser Asp Leu Tyr Pro Asp Ala Ser Thr Ala
145 150 155 160

Ser Asn Lys Ser Glu Gly Leu Ala Val Leu Ala Val Leu Ile Glu Met
165 170 175

Gly Ser Phe Asn Pro Ser Tyr Asp Lys Ile Phe Ser His Leu Gln His
180 185 190

Val Lys Tyr Lys Gly Gln Glu Ala Phe Val Pro Gly Phe Asn Ile Glu
195 200 205

Glu Leu Leu Pro Glu Arg Thr Ala Glu Tyr Tyr Arg Tyr Arg Gly Ser
210 215 220

Leu Thr Thr Pro Pro Cys Asn Pro Thr Val Leu Trp Thr Val Phe Arg
225 230 235 240

Asn Pro Val Gln Ile Ser Gln Glu Gln Leu Leu Ala Leu Glu Thr Ala
245 250 255

Leu Tyr Cys Thr His Met Asp Asp Pro Ser Pro Arg Glu Met Ile Asn
260 265 270

Asn Phe Arg Gln Val Gln Lys Phe Asp Glu Arg Leu Val Tyr Thr Ser
 275 280 285

Phe Ser Gln Val Gln Val Cys Thr Ala Ala Gly Leu Ser Leu Gly Ile
 290 295 300

Ile Leu Ser Leu Ala Leu Ala Gly Ile Leu Gly Ile Cys Ile Val Val
 305 310 315 320

Val Val Ser Ile Trp Leu Phe Arg Arg Lys Ser Ile Lys Lys Gly Asp
 325 330 335

Asn Lys Gly Val Ile Tyr Lys Pro Ala Thr Lys Met Glu Thr Glu Ala
 340 345 350

His Ala

<210> 217
 <211> 244
 <212> PRT
 <213> Homo sapiens

<400> 217

Met Glu Leu Phe Leu Ala Gly Arg Arg Val Leu Val Thr Gly Ala Gly
 1 5 10 15

Lys Gly Ile Gly Arg Gly Thr Val Gln Ala Leu His Ala Thr Gly Ala
 20 25 30

Arg Val Val Ala Val Ser Arg Thr Gln Ala Asp Leu Asp Ser Leu Val
 35 40 45

Arg Glu Cys Pro Gly Ile Glu Pro Val Cys Val Asp Leu Gly Asp Trp
 50 55 60

Glu Ala Thr Glu Arg Ala Leu Gly Ser Val Gly Pro Val Asp Leu Leu
 65 70 75 80

Val Asn Asn Ala Ala Val Ala Leu Leu Gln Pro Phe Leu Glu Val Thr
 85 90 95

Lys Glu Ala Phe Asp Arg Ser Phe Glu Val Asn Leu Arg Ala Val Ile

100 105 110
 Gln Val Ser Gln Ile Val Ala Arg Gly Leu Ile Ala Arg Gly Val Pro
 115 120 125
 Gly Ala Ile Val Asn Val Ser Ser Gln Cys Ser Gln Arg Ala Val Thr
 130 135 140
 Asn His Ser Val Tyr Cys Ser Thr Lys Gly Ala Leu Asp Met Leu Thr
 145 150 155 160
 Lys Val Met Ala Leu Glu Leu Gly Pro His Lys Ile Arg Val Asn Ala
 165 170 175
 Val Asn Pro Thr Val Val Met Thr Ser Met Gly Gln Ala Thr Trp Ser
 180 185 190
 Asp Pro His Lys Ala Lys Thr Met Leu Asn Arg Ile Pro Leu Gly Lys
 195 200 205
 Phe Ala Glu Val Glu His Val Val Asn Ala Ile Leu Phe Leu Leu Ser
 210 215 220
 Asp Arg Ser Gly Met Thr Thr Gly Ser Thr Leu Pro Val Glu Gly Gly
 225 230 235 240
 Phe Trp Ala Cys

<210> 218
 <211> 756
 <212> PRT
 <213> Homo sapiens

<400> 218

Met Ala Glu Ala His Gln Ala Val Ala Phe Gln Phe Thr Val Thr Pro
 1 5 10 15
 Asp Gly Ile Asp Leu Arg Leu Ser His Glu Ala Leu Arg Gln Ile Tyr
 20 25 30
 Leu Ser Gly Leu His Ser Trp Lys Lys Lys Phe Ile Arg Phe Lys Asn
 35 40 45

Gly Ile Ile Thr Gly Val Tyr Pro Ala Ser Pro Ser Ser Trp Leu Ile
 50 55 60

Val Val Val Gly Val Met Thr Thr Met Tyr Ala Lys Ile Asp Pro Ser
 65 70 75 80

Leu Gly Ile Ile Ala Lys Ile Asn Arg Thr Leu Glu Thr Ala Asn Cys
 85 90 95

Met Ser Ser Gln Thr Lys Asn Val Val Ser Gly Val Leu Phe Gly Thr
 100 105 110

Gly Leu Trp Val Ala Leu Ile Val Thr Met Arg Tyr Ser Leu Lys Val
 115 120 125

Leu Leu Ser Tyr His Gly Trp Met Phe Thr Glu His Gly Lys Met Ser
 130 135 140

Arg Ala Thr Lys Ile Trp Met Gly Met Val Lys Ile Phe Ser Gly Arg
 145 150 155 160

Lys Pro Met Leu Tyr Ser Phe Gln Thr Ser Leu Pro Arg Leu Pro Val
 165 170 175

Pro Ala Val Lys Asp Thr Val Asn Arg Tyr Leu Gln Ser Val Arg Pro
 180 185 190

Leu Met Lys Glu Glu Asp Phe Lys Arg Met Thr Ala Leu Ala Gln Asp
 195 200 205

Phe Ala Val Gly Leu Gly Pro Arg Leu Gln Trp Tyr Leu Lys Leu Lys
 210 215 220

Ser Trp Trp Ala Thr Asn Tyr Val Ser Asp Trp Trp Glu Glu Tyr Ile
 225 230 235 240

Tyr Leu Arg Gly Arg Gly Pro Leu Met Val Asn Ser Asn Tyr Tyr Ala
 245 250 255

Met Asp Leu Leu Tyr Ile Leu Pro Thr His Ile Gln Ala Ala Arg Ala
 260 265 270

Gly Asn Ala Ile His Ala Ile Leu Leu Tyr Arg Arg Lys Leu Asp Arg
 275 280 285

Glu Glu Ile Lys Pro Ile Arg Leu Leu Gly Ser Thr Ile Pro Leu Cys
 290 295 300

Ser Ala Gln Trp Glu Arg Met Phe Asn Thr Ser Arg Ile Pro Gly Glu
 305 310 315 320

Glu Thr Asp Thr Ile Gln His Met Arg Asp Ser Lys His Ile Val Val
 325 330 335

Tyr His Arg Gly Arg Tyr Phe Lys Val Trp Leu Tyr His Asp Gly Arg
 340 345 350

Leu Leu Lys Pro Arg Glu Met Glu Gln Gln Met Gln Arg Ile Leu Asp
 355 360 365

Asn Thr Ser Glu Pro Gln Pro Gly Glu Ala Arg Leu Ala Ala Leu Thr
 370 375 380

Ala Gly Asp Arg Val Pro Trp Ala Arg Cys Arg Gln Ala Tyr Phe Gly
 385 390 395 400

Arg Gly Lys Asn Lys Gln Ser Leu Asp Ala Val Glu Lys Ala Ala Phe
 405 410 415

Phe Val Thr Leu Asp Glu Thr Glu Glu Gly Tyr Arg Ser Glu Asp Pro
 420 425 430

Asp Thr Ser Met Asp Ser Tyr Ala Lys Ser Leu Leu His Gly Arg Cys
 435 440 445

Tyr Asp Arg Trp Phe Asp Lys Ser Phe Thr Phe Val Val Phe Lys Asn
 450 455 460

Gly Lys Met Gly Leu Asn Ala Glu His Ser Trp Ala Asp Ala Pro Ile
 465 470 475 480

Val Ala His Leu Trp Glu Tyr Val Met Ser Ile Asp Ser Leu Gln Leu
 485 490 495

Gly Tyr Ala Glu Asp Gly His Cys Lys Gly Asp Ile Asn Pro Asn Ile

500	505	510
Pro Tyr Pro Thr Arg Leu Gln Trp Asp Ile Pro Gly Glu Cys Gln Glu		
515	520	525
Val Ile Glu Thr Ser Leu Asn Thr Ala Asn Leu Leu Ala Asn Asp Val		
530	535	540
Asp Phe His Ser Phe Pro Phe Val Ala Phe Gly Lys Gly Ile Ile Lys		
545	550	555
Lys Cys Arg Thr Ser Pro Asp Ala Phe Val Gln Leu Ala Leu Gln Leu		
565	570	575
Ala His Tyr Lys Asp Met Gly Lys Phe Cys Leu Thr Tyr Glu Ala Ser		
580	585	590
Met Thr Arg Leu Phe Arg Glu Gly Arg Thr Glu Thr Val Arg Ser Cys		
595	600	605
Thr Thr Glu Ser Cys Asp Phe Val Arg Ala Met Val Asp Pro Ala Gln		
610	615	620
Thr Val Glu Gln Arg Leu Lys Leu Phe Lys Leu Ala Ser Glu Lys His		
625	630	635
Gln His Met Tyr Arg Leu Ala Met Thr Gly Ser Gly Ile Asp Arg His		
645	650	655
Leu Phe Cys Leu Tyr Val Val Ser Lys Tyr Leu Ala Val Glu Ser Pro		
660	665	670
Phe Leu Lys Glu Val Leu Ser Glu Pro Trp Arg Leu Ser Thr Ser Gln		
675	680	685
Thr Pro Gln Gln Gln Val Glu Leu Phe Asp Leu Glu Asn Asn Pro Glu		
690	695	700
Tyr Val Ser Ser Gly Gly Gly Phe Gly Pro Val Ala Asp Asp Gly Tyr		
705	710	715
Gly Val Ser Tyr Ile Leu Val Gly Glu Asn Leu Ile Asn Phe His Ile		
725	730	735

Ser Ser Lys Phe Ser Cys Pro Glu Thr Gly Ile Ile Ser Gln Gly Pro
 740 745 750

Ser Ser Asp Thr
 755

<210> 219
 <211> 509
 <212> PRT
 <213> Homo sapiens

<400> 219

Met Gly Cys Ser Ala Lys Ala Arg Trp Ala Ala Gly Ala Leu Gly Val
 1 5 10 15

Ala Gly Leu Leu Cys Ala Val Leu Gly Ala Val Met Ile Val Met Val
 20 25 30

Pro Ser Leu Ile Lys Gln Gln Val Leu Lys Asn Val Arg Ile Asp Pro
 35 40 45

Ser Ser Leu Ser Phe Asn Met Trp Lys Glu Ile Pro Ile Pro Phe Tyr
 50 55 60

Leu Ser Val Tyr Phe Phe Asp Val Met Asn Pro Ser Glu Ile Leu Lys
 65 70 75 80

Gly Glu Lys Pro Gln Val Arg Glu Arg Gly Pro Tyr Val Tyr Arg Glu
 85 90 95

Ser Arg His Lys Ser Asn Ile Thr Phe Asn Asn Asn Asp Thr Val Ser
 100 105 110

Phe Leu Glu Tyr Arg Thr Phe Gln Phe Gln Pro Ser Lys Ser His Gly
 115 120 125

Ser Glu Ser Asp Tyr Ile Val Met Pro Asn Ile Leu Val Leu Gly Ala
 130 135 140

Ala Val Met Met Glu Asn Lys Pro Met Thr Leu Lys Leu Ile Met Thr
 145 150 155 160

Leu Ala Phe Thr Thr Leu Gly Glu Arg Ala Phe Met Asn Arg Thr Val
 165 170 175

Gly Glu Ile Met Trp Gly Tyr Lys Asp Pro Leu Val Asn Leu Ile Asn
 180 185 190

Lys Tyr Phe Pro Gly Met Phe Pro Phe Lys Asp Lys Phe Gly Leu Phe
 195 200 205

Ala Glu Leu Asn Asn Ser Asp Ser Gly Leu Phe Thr Val Phe Thr Gly
 210 215 220

Val Gln Asn Ile Ser Arg Ile His Leu Val Asp Lys Trp Asn Gly Leu
 225 230 235 240

Ser Lys Val Asp Phe Trp His Ser Asp Gln Cys Asn Met Ile Asn Gly
 245 250 255

Thr Ser Gly Gln Met Trp Pro Pro Phe Met Thr Pro Glu Ser Ser Leu
 260 265 270

Glu Phe Tyr Ser Pro Glu Ala Cys Arg Ser Met Lys Leu Met Tyr Lys
 275 280 285

Glu Ser Gly Val Phe Glu Gly Ile Pro Thr Tyr Arg Phe Val Ala Pro
 290 295 300

Lys Thr Leu Phe Ala Asn Gly Ser Ile Tyr Pro Pro Asn Glu Gly Phe
 305 310 315 320

Cys Pro Cys Leu Glu Ser Gly Ile Gln Asn Val Ser Thr Cys Arg Phe
 325 330 335

Ser Ala Pro Leu Phe Leu Ser His Pro His Phe Leu Asn Ala Asp Pro
 340 345 350

Val Leu Ala Glu Ala Val Thr Gly Leu His Pro Asn Gln Glu Ala His
 355 360 365

Ser Leu Phe Leu Asp Ile His Pro Val Thr Gly Ile Pro Met Asn Cys
 370 375 380

Ser Val Lys Leu Gln Leu Ser Leu Tyr Met Lys Ser Val Ala Gly Ile

Leu Gln Gly Ala Arg Ile Leu Gly Ile Pro Val Ile Val Thr Glu Gln
85 90 95

Tyr Pro Lys Gly Leu Gly Ser Thr Val Gln Glu Ile Asp Leu Thr Gly
100 105 110

Val Lys Leu Val Leu Pro Lys Thr Lys Phe Ser Met Val Leu Pro Glu
115 120 125

Val Glu Ala Ala Leu Ala Glu Ile Pro Gly Val Arg Ser Val Val Leu
130 135 140

Phe Gly Val Glu Thr His Val Cys Ile Gln Gln Thr Ala Leu Glu Leu
145 150 155 160

Val Gly Arg Gly Val Glu Val His Ile Val Ala Asp Ala Thr Ser Ser
165 170 175

Arg Ser Met Met Asp Arg Met Phe Ala Arg Leu Thr Ser Arg Ser Asn
180 185 190

Gly Asp His Ser Asp His Glu
195

<210> 221
<211> 283
<212> PRT
<213> Homo sapiens

<400> 221

Met Thr Ser Gly Pro Gly Gly Pro Ala Ala Ala Ala Gly Gly Arg Lys
1 5 10 15

Glu Asn His Gln Trp Tyr Val Cys Asn Arg Glu Lys Leu Cys Glu Ser
20 25 30

Leu Gln Ala Val Phe Val Gln Ser Tyr Leu Asp Gln Gly Thr Gln Ile
35 40 45

Phe Leu Asn Asn Ser Ile Glu Lys Ser Gly Trp Leu Phe Ile Gln Leu
50 55 60

Tyr His Ser Phe Val Ser Ser Val Phe Ser Leu Phe Met Ser Arg Thr

65	70	75	80
Ser Ile Asn Gly Leu Leu Gly Arg Gly Ser Met Phe Val Phe Ser Pro	85	90	95
Asp Gln Phe Gln Arg Leu Leu Lys Ile Asn Pro Asp Trp Lys Thr His	100	105	110
Arg Leu Leu Asp Leu Gly Ala Gly Asp Gly Glu Val Thr Lys Ile Met	115	120	125
Ser Pro His Phe Glu Glu Ile Tyr Ala Thr Glu Leu Ser Glu Thr Met	130	135	140
Ile Trp Gln Leu Gln Lys Lys Lys Tyr Arg Val Leu Gly Ile Asn Glu	145	150	155
Trp Gln Asn Thr Gly Phe Gln Tyr Asp Val Ile Ser Cys Leu Asn Leu	165	170	175
Leu Asp Arg Cys Asp Gln Pro Leu Thr Leu Leu Lys Asp Ile Arg Ser	180	185	190
Val Leu Glu Pro Thr Arg Gly Arg Val Ile Leu Ala Leu Val Leu Pro	195	200	205
Phe His Pro Tyr Val Glu Asn Val Gly Gly Lys Trp Glu Lys Pro Ser	210	215	220
Glu Ile Leu Glu Ile Lys Gly Gln Asn Trp Glu Glu Gln Val Asn Ser	225	230	235
Leu Pro Glu Val Phe Arg Lys Ala Gly Phe Val Ile Glu Ala Phe Thr	245	250	255
Arg Leu Pro Tyr Leu Cys Glu Gly Asp Met Tyr Asn Asp Tyr Tyr Val	260	265	270
Leu Asp Asp Ala Val Phe Val Leu Lys Pro Val	275	280	
<210> 222			
<211> 220			

<212> PRT

<213> Homo sapiens

<400> 222

Met Ser Met Gly Leu Glu Ile Thr Gly Thr Ala Leu Ala Val Leu Gly
 1 5 10 15

Trp Leu Gly Thr Ile Val Cys Cys Ala Leu Pro Met Trp Arg Val Ser
 20 25 30

Ala Phe Ile Gly Ser Asn Ile Ile Thr Ser Gln Asn Ile Trp Glu Gly
 35 40 45

Leu Trp Met Asn Cys Val Val Gln Ser Thr Gly Gln Met Gln Cys Lys
 50 55 60

Val Tyr Asp Ser Leu Leu Ala Leu Pro Gln Asp Leu Gln Ala Ala Arg
 65 70 75 80

Ala Leu Ile Val Val Ala Ile Leu Leu Ala Ala Phe Gly Leu Leu Val
 85 90 95

Ala Leu Val Gly Ala Gln Cys Thr Asn Cys Val Gln Asp Asp Thr Ala
 100 105 110

Lys Ala Lys Ile Thr Ile Val Ala Gly Val Leu Phe Leu Leu Ala Ala
 115 120 125

Leu Leu Thr Leu Val Pro Val Ser Trp Ser Ala Asn Thr Ile Ile Arg
 130 135 140

Asp Phe Tyr Asn Pro Val Val Pro Glu Ala Gln Lys Arg Glu Met Gly
 145 150 155 160

Ala Gly Leu Tyr Val Gly Trp Ala Ala Ala Leu Gln Leu Leu Gly
 165 170 175

Gly Ala Leu Leu Cys Cys Ser Cys Pro Pro Arg Glu Lys Lys Tyr Thr
 180 185 190

Ala Thr Lys Val Val Tyr Ser Ala Pro Arg Ser Thr Gly Pro Gly Ala
 195 200 205

Ser Leu Gly Thr Gly Tyr Asp Arg Lys Asp Tyr Val
 210 215 220

<210> 223
 <211> 251
 <212> PRT
 <213> Homo sapiens
 <400> 223

Met Glu Gly Gly Ala Ala Ala Ala Thr Pro Thr Ala Leu Pro Tyr Tyr
 1 5 10 15

Val Ala Phe Ser Gln Leu Leu Gly Leu Thr Leu Val Ala Met Thr Gly
 20 25 30

Ala Trp Leu Gly Leu Tyr Arg Gly Gly Ile Ala Trp Glu Ser Asp Leu
 35 40 45

Gln Phe Asn Ala His Pro Leu Cys Met Val Ile Gly Leu Ile Phe Leu
 50 55 60

Gln Gly Asn Ala Leu Leu Val Tyr Arg Val Phe Arg Asn Glu Ala Lys
 65 70 75 80

Arg Thr Thr Lys Val Leu His Gly Leu Leu His Ile Phe Ala Leu Val
 85 90 95

Ile Ala Leu Val Gly Leu Val Ala Val Phe Asp Tyr His Arg Lys Lys
 100 105 110

Gly Tyr Ala Asp Leu Tyr Ser Leu His Ser Trp Cys Gly Ile Leu Val
 115 120 125

Phe Val Leu Tyr Phe Val Gln Trp Leu Val Gly Phe Ser Phe Phe Leu
 130 135 140

Phe Pro Gly Ala Ser Phe Ser Leu Arg Ser Arg Tyr Arg Pro Gln His
 145 150 155 160

Ile Phe Phe Gly Ala Thr Ile Phe Leu Leu Pro Val Gly Thr Ala Leu
 165 170 175

Leu Gly Leu Lys Glu Ala Leu Leu Phe Asn Leu Gly Gly Lys Tyr Ser
 180 185 190

Ala Phe Glu Pro Glu Gly Val Leu Ala Asn Val Leu Gly Leu Leu Leu
 195 200 205

Ala Cys Phe Gly Gly Ala Val Leu Tyr Ile Leu Thr Arg Ala Asp Trp
 210 215 220

Lys Arg Pro Ser Gln Ala Glu Glu Gln Ala Leu Ser Met Asp Phe Lys
 225 230 235 240

Thr Leu Arg Gln Gly Asp Ser Pro Gly Ser Gln
 245 250

<210> 224
 <211> 401
 <212> PRT
 <213> Homo sapiens

<400> 224

Tyr Val Cys Asn Cys Ser Val Val Gly Ser Leu Asn Val Asn Arg Cys
 1 5 10 15

Asn Gln Thr Thr Gly Gln Cys Glu Cys Arg Pro Gly Tyr Gln Gly Leu
 20 25 30

His Cys Glu Thr Cys Lys Glu Gly Phe Tyr Leu Asn Tyr Thr Ser Gly
 35 40 45

Leu Cys Gln Pro Cys Asp Cys Ser Pro His Gly Ala Leu Ser Ile Pro
 50 55 60

Cys Asn Ser Ser Gly Lys Cys Gln Cys Lys Val Gly Val Ile Gly Ser
 65 70 75 80

Ile Cys Asp Arg Cys Gln Asp Gly Tyr Tyr Gly Phe Ser Lys Asn Gly
 85 90 95

Cys Leu Pro Cys Gln Cys Asn Asn Arg Ser Ala Ser Cys Asp Ala Leu
 100 105 110

Thr Gly Ala Cys Leu Asn Cys Gln Glu Asn Ser Lys Gly Asn His Cys
 115 120 125

Glu Glu Cys Lys Glu Gly Phe Tyr Gln Ser Pro Asp Ala Thr Lys Glu
 130 135 140

Cys Leu Arg Cys Pro Cys Ser Ala Val Thr Ser Thr Gly Ser Cys Ser
 145 150 155 160

Ile Lys Ser Ser Glu Leu Glu Pro Glu Cys Asp Gln Cys Lys Asp Gly
 165 170 175

Tyr Ile Gly Pro Asn Cys Asn Lys Cys Glu Asn Gly Tyr Tyr Asn Phe
 180 185 190

Asp Ser Ile Cys Arg Lys Cys Gln Cys His Gly His Val Asp Pro Val
 195 200 205

Lys Thr Pro Lys Ile Cys Lys Pro Glu Ser Gly Glu Cys Ile Asn Cys
 210 215 220

Leu His Asn Thr Thr Gly Phe Trp Cys Glu Asn Cys Leu Glu Gly Tyr
 225 230 235 240

Val His Asp Leu Glu Gly Asn Cys Ile Lys Lys Glu Val Ile Leu Pro
 245 250 255

Thr Pro Glu Gly Ser Thr Ile Leu Val Ser Asn Ala Ser Leu Thr Thr
 260 265 270

Ser Val Pro Thr Pro Val Ile Asn Ser Thr Phe Thr Pro Thr Thr Leu
 275 280 285

Gln Thr Ile Phe Ser Val Ser Thr Ser Glu Asn Ser Thr Ser Ala Leu
 290 295 300

Ala Asp Val Ser Trp Thr Gln Phe Asn Ile Ile Ile Leu Thr Val Ile
 305 310 315 320

Ile Ile Val Val Val Leu Leu Met Gly Phe Val Gly Ala Val Tyr Met
 325 330 335

Tyr Arg Glu Tyr Gln Asn Arg Lys Leu Asn Ala Pro Phe Trp Thr Ile
 340 345 350

Glu Leu Lys Glu Asp Asn Ile Ser Phe Ser Ser Tyr His Asp Ser Ile

355

360

365

Pro Asn Ala Asp Val Ser Gly Leu Leu Glu Asp Asp Gly Asn Glu Val
 370 375 380

Ala Pro Asn Gly Gln Leu Thr Leu Thr Thr Pro Ile His Asn Tyr Lys
 385 390 395 400

Ala

<210> 225
 <211> 686
 <212> PRT
 <213> Homo sapiens

<400> 225

Met Lys Pro Ser Trp Leu Gln Cys Arg Lys Val Thr Ser Ala Gly Gly
 1 5 10 15

Leu Gly Gly Pro Leu Pro Gly Ser Ser Pro Ala Arg Gly Ala Gly Ala
 20 25 30

Ala Leu Arg Ala Leu Val Val Pro Gly Pro Arg Gly Gly Leu Gly Gly
 35 40 45

Arg Gly Cys Arg Ala Leu Ser Ser Gly Ser Gly Ser Glu Tyr Lys Thr
 50 55 60

His Phe Ala Ala Ser Val Thr Asp Pro Glu Arg Phe Trp Gly Lys Ala
 65 70 75 80

Ala Glu Gln Ile Ser Trp Tyr Lys Pro Trp Thr Lys Thr Leu Glu Asn
 85 90 95

Lys His Ser Pro Ser Thr Arg Trp Phe Val Glu Gly Met Leu Asn Ile
 100 105 110

Cys Tyr Asn Ala Val Asp Arg His Ile Glu Asn Gly Lys Gly Asp Lys
 115 120 125

Ile Ala Ile Ile Tyr Asp Ser Pro Val Thr Asn Thr Lys Ala Thr Phe
 130 135 140

Thr Tyr Lys Glu Val Leu Glu Gln Val Ser Lys Leu Ala Gly Val Leu
 145 150 155 160

Val Lys His Gly Ile Lys Lys Gly Asp Thr Val Val Ile Tyr Met Pro
 165 170 175

Met Ile Pro Gln Ala Met Tyr Thr Met Leu Ala Cys Ala Arg Ile Gly
 180 185 190

Ala Ile His Ser Leu Ile Phe Gly Gly Phe Ala Ser Lys Glu Leu Ser
 195 200 205

Ser Arg Ile Asp His Val Lys Pro Lys Val Val Val Thr Ala Ser Phe
 210 215 220

Gly Ile Glu Pro Gly Arg Arg Val Glu Tyr Val Pro Leu Val Glu Glu
 225 230 235 240

Ala Leu Lys Ile Gly Gln His Lys Pro Asp Lys Ile Leu Ile Tyr Asn
 245 250 255

Arg Pro Asn Met Glu Ala Val Pro Leu Ala Pro Gly Arg Asp Leu Asp
 260 265 270

Trp Asp Glu Glu Met Ala Lys Ala Gln Ser His Asp Cys Val Pro Val
 275 280 285

Leu Ser Glu His Pro Leu Tyr Ile Leu Tyr Thr Ser Gly Thr Thr Gly
 290 295 300

Leu Pro Lys Gly Val Ile Arg Pro Thr Gly Gly Tyr Ala Val Met Leu
 305 310 315 320

His Trp Ser Met Ser Ser Ile Tyr Gly Leu Gln Pro Gly Glu Val Trp
 325 330 335

Trp Ala Ala Ser Asp Leu Gly Trp Val Val Gly His Ser Tyr Ile Cys
 340 345 350

Tyr Gly Pro Leu Leu His Gly Asn Thr Thr Val Leu Tyr Glu Gly Lys
 355 360 365

Pro Val Gly Thr Pro Asp Ala Gly Ala Tyr Phe Arg Val Leu Ala Glu
 370 375 380

His Gly Val Ala Ala Leu Phe Thr Ala Pro Thr Ala Ile Arg Ala Ile
 385 390 395 400

Arg Gln Gln Asp Pro Gly Ala Ala Leu Gly Lys Gln Tyr Ser Leu Thr
 405 410 415

Arg Phe Lys Thr Leu Phe Val Ala Gly Glu Arg Cys Asp Val Glu Thr
 420 425 430

Leu Glu Trp Ser Lys Asn Val Phe Arg Val Pro Val Leu Asp His Trp
 435 440 445

Trp Gln Thr Glu Thr Gly Ser Pro Ile Thr Ala Ser Cys Val Gly Leu
 450 455 460

Gly Asn Ser Lys Thr Pro Pro Pro Gly Gln Ala Gly Lys Ser Val Pro
 465 470 475 480

Gly Tyr Asn Val Met Ile Leu Asp Asp Asn Met Gln Lys Leu Lys Ala
 485 490 495

Arg Cys Leu Gly Asn Ile Val Val Lys Leu Pro Leu Pro Pro Gly Ala
 500 505 510

Phe Ser Gly Leu Trp Lys Asn Gln Glu Ala Phe Lys His Leu Tyr Phe
 515 520 525

Glu Lys Phe Pro Gly Tyr Tyr Asp Thr Met Asp Ala Gly Tyr Met Asp
 530 535 540

Glu Glu Gly Tyr Leu Tyr Val Met Ser Arg Val Asp Asp Val Ile Asn
 545 550 555 560

Val Ala Gly His Arg Ile Ser Ala Gly Ala Ile Glu Glu Ser Ile Leu
 565 570 575

Ser His Gly Thr Val Ala Asp Cys Ala Val Val Gly Lys Glu Asp Pro
 580 585 590

Leu Lys Gly His Val Pro Leu Ala Leu Cys Val Leu Arg Lys Asp Ile

595 600 605
 Asn Ala Thr Glu Glu Gln Val Leu Glu Glu Ile Val Lys His Val Arg
 610 615 620
 Gln Asn Ile Gly Pro Val Ala Ala Phe Arg Asn Ala Val Phe Val Lys
 625 630 635 640
 Gln Leu Pro Lys Thr Arg Ser Gly Lys Ile Pro Arg Ser Ala Leu Ser
 645 650 655
 Ala Ile Val Asn Gly Lys Pro Tyr Lys Ile Thr Ser Thr Ile Glu Asp
 660 665 670
 Pro Ser Ile Phe Gly His Val Glu Glu Met Leu Lys Gln Ala
 675 680 685

 <210> 226
 <211> 225
 <212> PRT
 <213> Homo sapiens

 <400> 226
 Met Ala Ala Ala Gly Gly Gly Gly Gly Gly Ala Ala Ala Ala Gly Arg
 1 5 10 15
 Ala Tyr Ser Phe Lys Val Val Leu Leu Gly Glu Gly Cys Val Gly Lys
 20 25 30
 Thr Ser Leu Val Leu Arg Tyr Cys Glu Asn Lys Phe Asn Asp Lys His
 35 40 45
 Ile Thr Thr Leu Gln Ala Ser Phe Leu Thr Lys Lys Leu Asn Ile Gly
 50 55 60
 Gly Lys Arg Val Asn Leu Ala Ile Trp Asp Thr Ala Gly Gln Glu Arg
 65 70 75 80
 Phe His Ala Leu Gly Pro Ile Tyr Tyr Arg Asp Ser Asn Gly Ala Ile
 85 90 95
 Leu Val Tyr Asp Ile Thr Asp Glu Asp Ser Phe Gln Lys Val Lys Asn
 100 105 110

Trp Val Lys Glu Leu Arg Lys Met Leu Gly Asn Glu Ile Cys Leu Cys
 115 120 125

Ile Val Gly Asn Lys Ile Asp Leu Glu Lys Glu Arg His Val Ser Ile
 130 135 140

Gln Glu Ala Glu Ser Tyr Ala Glu Ser Val Gly Ala Lys His Tyr His
 145 150 155 160

Thr Ser Ala Lys Gln Asn Lys Gly Ile Glu Glu Leu Phe Leu Asp Leu
 165 170 175

Cys Lys Arg Met Ile Glu Thr Ala Gln Val Asp Glu Arg Ala Lys Gly
 180 185 190

Asn Gly Ser Ser Gln Pro Gly Thr Ala Arg Arg Gly Val Gln Ile Ile
 195 200 205

Asp Asp Glu Pro Gln Ala Gln Thr Ser Gly Gly Gly Cys Cys Ser Ser
 210 215 220

Gly
 225

<210> 227
 <211> 380
 <212> PRT
 <213> Homo sapiens

<400> 227

Met Gly Ser Thr Asp Ser Lys Leu Asn Phe Arg Lys Ala Val Ile Gln
 1 5 10 15

Leu Thr Thr Lys Thr Gln Pro Val Glu Ala Thr Asp Asp Ala Phe Trp
 20 25 30

Asp Gln Phe Trp Ala Asp Thr Ala Thr Ser Val Gln Asp Val Phe Ala
 35 40 45

Leu Val Pro Ala Ala Glu Ile Arg Ala Val Arg Glu Glu Ser Pro Ser
 50 55 60

Asn Leu Ala Thr Leu Cys Tyr Lys Ala Val Glu Lys Leu Val Gln Gly

65 70 75 80
 Ala Glu Ser Gly Cys His Ser Glu Lys Glu Lys Gln Ile Val Leu Asn
 85 90 95
 Cys Ser Arg Leu Leu Thr Arg Val Leu Pro Tyr Ile Phe Glu Asp Pro
 100 105 110
 Asp Trp Arg Gly Phe Phe Trp Ser Thr Val Pro Gly Ala Gly Arg Gly
 115 120 125
 Gly Gly Glu Glu Asp Asp Glu His Ala Arg Pro Leu Ala Glu Ser Leu
 130 135 140
 Leu Leu Ala Ile Ala Asp Leu Leu Phe Cys Pro Asp Phe Thr Val Gln
 145 150 155 160
 Ser His Arg Arg Ser Thr Val Asp Ser Ala Glu Asp Val His Ser Leu
 165 170 175
 Asp Ser Cys Glu Tyr Ile Trp Glu Ala Gly Val Gly Phe Ala His Ser
 180 185 190
 Pro Gln Pro Asn Tyr Ile His Asp Met Asn Arg Met Glu Leu Leu Lys
 195 200 205
 Leu Leu Leu Thr Cys Phe Ser Glu Ala Met Tyr Leu Pro Pro Ala Pro
 210 215 220
 Glu Ser Gly Ser Thr Asn Pro Trp Val Gln Phe Phe Cys Ser Thr Glu
 225 230 235 240
 Asn Arg His Ala Leu Pro Leu Phe Thr Ser Leu Leu Asn Thr Val Cys
 245 250 255
 Ala Tyr Asp Pro Val Gly Tyr Gly Ile Pro Tyr Asn His Leu Leu Phe
 260 265 270
 Ser Asp Tyr Arg Glu Pro Leu Val Glu Glu Ala Ala Gln Val Leu Ile
 275 280 285
 Val Thr Leu Asp His Asp Ser Ala Ser Ser Ala Ser Pro Thr Val Asp
 290 295 300

Gly Thr Thr Thr Gly Thr Ala Met Asp Asp Ala Asp Pro Pro Gly Pro
305 310 315 320

Glu Asn Leu Phe Val Asn Tyr Leu Ser Arg Ile His Arg Glu Glu Asp
325 330 335

Phe Gln Phe Ile Leu Lys Gly Ile Ala Arg Leu Leu Ser Asn Leu Leu
340 345 350

Leu Gln Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys
355 360 365

Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys
370 375 380

<210> 228
<211> 144
<212> PRT
<213> Homo sapiens

<400> 228

Met Cys Arg Val Gln Thr His Gly Cys His Pro Leu Arg Ser Val Thr
1 5 10 15

Val Arg Pro Asp Pro Ser Pro Ala Ala Pro Pro Pro His Pro Gly Pro
20 25 30

Pro Arg Gln Leu Ser Gln Gly Ala Gln Ala Cys Leu Ala Pro Gln Pro
35 40 45

Ser Gly Asn Pro Ala Arg Arg Pro Leu Gln Val Gly Ser Gly Pro Gln
50 55 60

Val Ala Lys Gln Arg Gln Gln Gln Pro Arg Leu Thr Pro Cys Pro Ser
65 70 75 80

Leu Trp Arg Pro Gly Thr Pro Ala Ile Ser Thr Thr Trp Val Arg Leu
85 90 95

Ser Leu Ser Gly Ser Pro Ala Arg Val Pro Pro Gly Val Leu Gly His
100 105 110

Leu Arg Gly Ser Leu Met Gly Gln Ala Gly Gln Ser Glu Leu Arg Ala
 115 120 125

Leu Ser Gly Trp Cys Pro Asn Leu Ser Thr Pro His Ser Phe Pro Pro
 130 135 140

<210> 229
 <211> 141
 <212> PRT
 <213> Homo sapiens

<400> 229

Met Thr Lys Gln His Glu Leu Gly Gly Leu Leu Ala Leu Val Gln Asn
 1 5 10 15

Cys Gln Ser Glu Met Asn Ile Lys Asp Ser Arg Ala Val Gly Leu Ser
 20 25 30

Val Lys Arg Leu Cys Ile Ser Phe Val Asp Glu Phe Cys Glu Arg Thr
 35 40 45

Glu Arg Pro Leu Tyr Leu Ala Gln Gly Leu Phe Met Lys Arg Glu Thr
 50 55 60

Tyr Trp Glu Val Gln Asp Ser Gly Ile Ser Pro Leu Leu Leu Leu Leu
 65 70 75 80

Ser Thr Ala Leu Asp Cys Ser Pro Glu Ala Glu Thr Arg Gln Ser Pro
 85 90 95

Gly Gly Arg Lys Met Leu Gln Glu Pro Thr Leu Ser Met Ser Leu Gln
 100 105 110

Ile Leu Thr Gly Phe Leu Trp Val Gln Leu Trp Asn Trp Glu Thr Phe
 115 120 125

Leu Arg Ile Arg Thr His Ser Thr Asp Ala Ser Cys Pro
 130 135 140

<210> 230
 <211> 161
 <212> PRT
 <213> Homo sapiens

<400> 230

Met Ser Tyr Leu Ser Gly Ser Ser Ala Ser Pro Ala Arg Arg Leu Gly
1 5 10 15

Val Val Lys Val Val Pro Arg Pro Cys Leu Gln Trp Leu Glu Asn Pro
20 25 30

Gly Gly Cys Leu Gly Pro Ser Gly Gln Arg Glu Ala Gly Ser Ser Ser
35 40 45

Pro Gly Asp Cys Gly His Ile Gly Ala Cys Leu Gly Leu Glu Gly Gln
50 55 60

Val Thr Ser Pro Ala Thr Leu Pro Ser Leu Leu Trp Gly Pro His Phe
65 70 75 80

Arg Ala Thr Leu Pro Glu Ala His Ala Ser Ile His Ser Phe Ser Ala
85 90 95

Leu Asn Leu Ile His Lys Gln Pro Pro Pro Phe Pro Phe Pro Ser His
100 105 110

Ser Val Asp Val Ile Leu Pro Pro Pro Val Ser Ile Leu Arg Gln Ala
115 120 125

Ser Lys Glu Ala Leu Thr Leu Leu Pro Lys Trp Cys Phe Leu Lys Asn
130 135 140

Thr Ile Thr Thr Leu Gly Ala Ile Phe Ser His Leu Pro Val Phe Arg
145 150 155 160

Met

<210> 231
<211> 132
<212> PRT
<213> Homo sapiens

<400> 231

Met Arg Ala Ile Asn Ile Ala Asp Glu Leu Pro Arg Ser Arg Ala Arg
1 5 10 15

Lys Leu Ala Asp Glu Gln Leu Ser Ser Val Ile Gln Asp Met Ala Val

20 25 30
 Arg Gln His Leu Leu Thr Asn Leu Val Glu Val Asp Gly Arg Phe Val
 35 40 45
 Trp Arg Val Asn Leu Asp Ala Leu Thr Gln His Leu Asp Lys Ile Leu
 50 55 60
 Ala Phe Pro Gln Arg Gln Glu Ser Tyr Leu Gly Pro Thr Leu Phe Leu
 65 70 75 80
 Leu Gly Gly Asn Ser Gln Phe Val His Pro Ser His His Pro Glu Ile
 85 90 95
 Met Arg Leu Phe Pro Arg Ala Gln Met Gln Thr Val Pro Asn Ala Gly
 100 105 110
 His Trp Ile His Ala Asp Arg Pro Gln Asp Phe Ile Ala Ala Ile Arg
 115 120 125
 Gly Phe Leu Val
 130
 <210> 232
 <211> 328
 <212> PRT
 <213> Homo sapiens
 <400> 232
 Met Leu Pro Arg Val Gly Cys Pro Ala Leu Pro Leu Pro Pro Pro Pro
 1 5 10 15
 Leu Leu Pro Leu Leu Pro Leu Leu Leu Leu Leu Gly Ala Ser Gly
 20 25 30
 Gly Gly Gly Gly Ala Arg Ala Glu Val Leu Phe Arg Cys Pro Pro Cys
 35 40 45
 Thr Pro Glu Arg Leu Ala Ala Cys Gly Pro Pro Pro Val Ala Pro Pro
 50 55 60
 Ala Ala Val Ala Ala Val Ala Gly Gly Ala Arg Met Pro Cys Ala Glu
 65 70 75 80

Leu Val Arg Glu Pro Gly Cys Gly Cys Cys Ser Val Cys Ala Arg Leu
85 90 95

Glu Gly Glu Ala Cys Gly Val Tyr Thr Pro Arg Cys Gly Gln Gly Leu
100 105 110

Arg Cys Tyr Pro His Pro Gly Ser Glu Leu Pro Leu Gln Ala Leu Val
115 120 125

Met Gly Glu Gly Thr Cys Glu Lys Arg Arg Asp Ala Glu Tyr Gly Ala
130 135 140

Ser Pro Glu Gln Val Ala Asp Asn Gly Asp Asp His Ser Glu Gly Gly
145 150 155 160

Leu Val Glu Asn His Val Asp Ser Thr Met Asn Met Leu Gly Gly Gly
165 170 175

Gly Ser Ala Gly Arg Lys Pro Leu Lys Ser Gly Met Lys Glu Leu Ala
180 185 190

Val Phe Arg Glu Lys Val Thr Glu Gln His Arg Gln Met Gly Lys Gly
195 200 205

Gly Lys His His Leu Gly Leu Glu Glu Pro Lys Lys Leu Arg Pro Pro
210 215 220

Pro Ala Arg Thr Pro Cys Gln Gln Glu Leu Asp Gln Val Leu Glu Arg
225 230 235 240

Ile Ser Thr Met Arg Leu Pro Asp Glu Arg Gly Pro Leu Glu His Leu
245 250 255

Tyr Ser Leu His Ile Pro Asn Cys Asp Lys His Gly Leu Tyr Asn Leu
260 265 270

Lys Gln Cys Lys Met Ser Leu Asn Gly Gln Arg Gly Glu Cys Trp Cys
275 280 285

Val Asn Pro Asn Thr Gly Lys Leu Ile Gln Gly Ala Pro Thr Ile Arg
290 295 300

Gly Asp Pro Glu Cys His Leu Phe Tyr Asn Glu Gln Gln Glu Ala Cys
 305 310 315 320

Gly Val His Thr Gln Arg Met Gln
 325

<210> 233
 <211> 417
 <212> PRT
 <213> Homo sapiens

<400> 233

Met Ala Val Glu Thr Thr Val His Thr His Leu Ser Ala Ser Pro Pro
 1 5 10 15

Gln Gly Ser Pro Tyr Asp His Thr Pro Gly Met Ala Gly Ser Leu Gly
 20 25 30

Tyr His Pro Tyr Ala Ala Pro Leu Gly Ser Tyr Pro Tyr Gly Asp Pro
 35 40 45

Ala Tyr Arg Lys Asn Ala Thr Arg Asp Ala Thr Ala Thr Leu Lys Ala
 50 55 60

Trp Leu Asn Glu His Arg Lys Asn Pro Tyr Pro Thr Lys Gly Glu Lys
 65 70 75 80

Ile Met Leu Ala Ile Ile Thr Lys Met Thr Leu Thr Gln Val Ser Thr
 85 90 95

Trp Phe Ala Asn Ala Arg Arg Arg Leu Lys Lys Glu Asn Lys Met Thr
 100 105 110

Trp Thr Pro Arg Asn Arg Ser Glu Asp Glu Glu Glu Glu Glu Asn Ile
 115 120 125

Asp Leu Glu Lys Asn Asp Glu Asp Glu Pro Gln Lys Pro Glu Asp Lys
 130 135 140

Gly Asp Pro Glu Gly Pro Glu Ala Gly Gly Ala Glu Gln Lys Ala Ala
 145 150 155 160

Ser Gly Cys Glu Arg Leu Gln Gly Pro Pro Thr Pro Ala Gly Lys Glu
 165 170 175

Thr Glu Gly Ser Leu Ser Asp Ser Asp Phe Lys Glu Pro Pro Ser Glu
 180 185 190

Gly Arg Leu Asp Ala Leu Gln Gly Pro Pro Arg Thr Gly Gly Pro Ser
 195 200 205

Pro Ala Gly Pro Ala Ala Ala Arg Leu Ala Glu Asp Pro Ala Pro His
 210 215 220

Tyr Pro Ala Gly Ala Pro Ala Pro Gly Pro His Pro Ala Ala Gly Glu
 225 230 235 240

Val Pro Pro Gly Pro Gly Gly Pro Ser Val Ile His Ser Pro Pro Pro
 245 250 255

Pro Pro Pro Pro Ala Val Leu Ala Lys Pro Lys Leu Trp Ser Leu Ala
 260 265 270

Glu Ile Ala Thr Leu Ser Asp Lys Val Lys Asp Gly Gly Gly Gly Asn
 275 280 285

Glu Gly Ser Pro Cys Pro Pro Cys Pro Gly Pro Ile Ala Gly Gln Ala
 290 295 300

Leu Gly Gly Ser Arg Ala Ser Pro Ala Pro Ala Pro Ser Arg Ser Pro
 305 310 315 320

Ser Ala Gln Cys Pro Phe Pro Gly Gly Thr Val Leu Ser Arg Pro Leu
 325 330 335

Tyr Tyr Thr Ala Pro Phe Tyr Pro Gly Tyr Thr Asn Tyr Gly Ser Phe
 340 345 350

Gly His Leu His Gly His Pro Gly Pro Gly Pro Gly Pro Thr Thr Gly
 355 360 365

Pro Gly Ser His Phe Asn Gly Leu Asn Gln Thr Val Leu Asn Arg Ala
 370 375 380

Asp Ala Leu Ala Lys Asp Pro Lys Met Leu Arg Ser Gln Ser Gln Leu
 385 390 395 400

Asp Leu Cys Lys Asp Ser Pro Tyr Glu Leu Lys Lys Gly Met Ser Asp
 405 410 415

Ile

<210> 234
 <211> 257
 <212> PRT
 <213> Homo sapiens

<400> 234

Met Ser Gly His Lys Cys Ser Tyr Pro Trp Asp Leu Gln Asp Arg Tyr
 1 5 10 15

Ala Gln Asp Lys Ser Val Val Asn Lys Met Gln Gln Lys Tyr Trp Glu
 20 25 30

Thr Lys Gln Ala Phe Ile Lys Ala Thr Gly Lys Lys Glu Asp Glu His
 35 40 45

Val Val Ala Ser Asp Ala Asp Leu Asp Ala Lys Leu Glu Leu Phe His
 50 55 60

Ser Ile Gln Arg Thr Cys Leu Asp Leu Ser Lys Ala Ile Val Leu Tyr
 65 70 75 80

Gln Lys Arg Ile Cys Phe Leu Ser Gln Glu Glu Asn Glu Leu Gly Lys
 85 90 95

Phe Leu Arg Ser Gln Gly Phe Gln Asp Lys Thr Arg Ala Gly Lys Met
 100 105 110

Met Gln Ala Thr Gly Lys Ala Leu Cys Phe Ser Ser Gln Gln Arg Leu
 115 120 125

Ala Leu Arg Asn Pro Leu Cys Arg Phe His Gln Glu Val Glu Thr Phe
 130 135 140

Arg His Arg Ala Ile Ser Asp Thr Trp Leu Thr Val Asn Arg Met Glu
 145 150 155 160

Gln Cys Arg Thr Glu Tyr Arg Gly Ala Leu Leu Trp Met Lys Asp Val

165 170 175
 Ser Gln Glu Leu Asp Pro Asp Leu Tyr Lys Gln Met Glu Lys Phe Arg
 180 185 190
 Lys Val Gln Thr Gln Val Arg Leu Ala Lys Lys Asn Phe Asp Lys Leu
 195 200 205
 Lys Met Asp Val Cys Gln Lys Val Asp Leu Leu Gly Ala Ser Arg Cys
 210 215 220
 Asn Leu Leu Ser His Met Leu Ala Thr Tyr Gln Leu Ala Trp Asp Gln
 225 230 235 240
 Trp Gln Gly Pro Arg Asn Leu Lys Val Leu Thr Lys Met Thr Cys Cys
 245 250 255
 Cys
 <210> 235
 <211> 395
 <212> PRT
 <213> Homo sapiens
 <400> 235
 Met Asp Leu Gly Ile Pro Asp Leu Leu Asp Ala Trp Leu Glu Pro Pro
 1 5 10 15
 Glu Asp Ile Phe Ser Thr Gly Ser Val Leu Glu Leu Gly Leu His Cys
 20 25 30
 Pro Pro Leu Glu Val Pro Val Thr Arg Leu Gln Glu Gln Gly Leu Gln
 35 40 45
 Gly Trp Lys Ser Gly Gly Asp Arg Gly Cys Gly Leu Gln Glu Ser Glu
 50 55 60
 Pro Glu Asp Phe Leu Lys Leu Phe Ile Asp Pro Asn Glu Val Tyr Cys
 65 70 75 80
 Ser Glu Ala Ser Pro Gly Ser Asp Ser Gly Ile Ser Glu Asp Pro Cys
 85 90 95

His Pro Asp Ser Pro Pro Ala Pro Arg Ala Thr Ser Ser Pro Met Leu
 100 105 110

Tyr Glu Val Val Tyr Glu Ala Gly Ala Leu Glu Arg Met Gln Gly Glu
 115 120 125

Thr Gly Pro Asn Val Gly Leu Ile Ser Ile Gln Leu Asp Gln Trp Ser
 130 135 140

Pro Ala Phe Met Val Pro Asp Ser Cys Met Val Ser Glu Leu Pro Phe
 145 150 155 160

Asp Ala His Ala His Ile Leu Pro Arg Ala Gly Thr Val Ala Pro Val
 165 170 175

Pro Cys Thr Thr Leu Leu Pro Cys Gln Thr Leu Phe Leu Thr Asp Glu
 180 185 190

Glu Lys Arg Leu Leu Gly Gln Glu Gly Val Ser Leu Pro Ser His Leu
 195 200 205

Pro Leu Thr Lys Ala Glu Glu Arg Val Leu Lys Lys Val Arg Arg Lys
 210 215 220

Ile Arg Asn Lys Gln Ser Ala Gln Asp Ser Arg Arg Arg Lys Lys Glu
 225 230 235 240

Tyr Ile Asp Gly Leu Glu Ser Arg Val Ala Ala Cys Ser Ala Gln Asn
 245 250 255

Gln Glu Leu Gln Lys Lys Val Gln Glu Leu Glu Arg His Asn Ile Ser
 260 265 270

Leu Val Ala Gln Leu Arg Gln Leu Gln Thr Leu Ile Ala Gln Thr Ser
 275 280 285

Asn Lys Ala Ala Gln Thr Ser Thr Cys Val Leu Ile Leu Leu Phe Ser
 290 295 300

Leu Ala Leu Ile Ile Leu Pro Ser Phe Ser Pro Phe Gln Ser Arg Pro
 305 310 315 320

Glu Ala Gly Ser Glu Asp Tyr Gln Pro His Gly Val Thr Ser Arg Asn
 325 330 335

Ile Leu Thr His Lys Asp Val Thr Glu Asn Leu Glu Thr Gln Val Val
 340 345 350

Glu Ser Arg Leu Arg Glu Pro Pro Gly Ala Lys Asp Ala Asn Gly Ser
 355 360 365

Thr Arg Thr Leu Leu Glu Lys Met Gly Gly Lys Pro Arg Pro Ser Gly
 370 375 380

Arg Ile Arg Ser Val Leu His Ala Asp Glu Met
 385 390 395

<210> 236
 <211> 351
 <212> PRT
 <213> Homo sapiens

<400> 236

Met Ala Ala Ala Pro Leu Lys Val Cys Ile Val Gly Ser Gly Asn Trp
 1 5 10 15

Gly Ser Ala Val Ala Lys Ile Ile Gly Asn Asn Val Lys Lys Leu Gln
 20 25 30

Lys Phe Ala Ser Thr Val Lys Met Trp Val Phe Glu Glu Thr Val Asn
 35 40 45

Gly Arg Lys Leu Thr Asp Ile Ile Asn Asn Asp His Glu Asn Val Lys
 50 55 60

Tyr Leu Pro Gly His Lys Leu Pro Glu Asn Val Val Ala Met Ser Asn
 65 70 75 80

Leu Ser Glu Ala Val Gln Asp Ala Asp Leu Leu Val Phe Val Ile Pro
 85 90 95

His Gln Phe Ile His Arg Ile Cys Asp Glu Ile Thr Gly Arg Val Pro
 100 105 110

Lys Lys Ala Leu Gly Ile Thr Leu Ile Lys Gly Ile Asp Glu Gly Pro
 115 120 125

Glu Gly Leu Lys Leu Ile Ser Asp Ile Ile Arg Glu Lys Met Gly Ile
 130 135 140

Asp Ile Ser Val Leu Met Gly Ala Asn Ile Ala Asn Glu Val Ala Ala
 145 150 155 160

Glu Lys Phe Cys Glu Thr Thr Ile Gly Ser Lys Val Met Glu Asn Gly
 165 170 175

Leu Leu Phe Lys Glu Leu Leu Gln Thr Pro Asn Phe Arg Ile Thr Val
 180 185 190

Val Asp Asp Ala Asp Thr Val Glu Leu Cys Gly Ala Leu Lys Asn Ile
 195 200 205

Val Ala Val Gly Ala Gly Phe Cys Asp Gly Leu Arg Cys Gly Asp Asn
 210 215 220

Thr Lys Ala Ala Val Ile Arg Leu Gly Leu Met Glu Met Ile Ala Phe
 225 230 235 240

Ala Arg Ile Phe Cys Lys Gly Gln Val Ser Thr Ala Thr Phe Leu Glu
 245 250 255

Ser Cys Gly Val Ala Asp Leu Ile Thr Thr Cys Tyr Gly Gly Arg Asn
 260 265 270

Arg Arg Val Ala Glu Ala Phe Ala Arg Thr Gly Lys Thr Ile Glu Glu
 275 280 285

Leu Glu Lys Glu Met Leu Asn Gly Gln Lys Leu Gln Gly Pro Gln Thr
 290 295 300

Ser Ala Glu Val Tyr Arg Ile Leu Lys Gln Lys Gly Leu Leu Asp Lys
 305 310 315 320

Phe Pro Leu Phe Thr Ala Val Tyr Gln Ile Cys Tyr Glu Ser Arg Pro
 325 330 335

Val Gln Glu Met Leu Ser Cys Leu Gln Ser His Pro Glu His Thr
 340 345 350

<210> 237
 <211> 871
 <212> PRT
 <213> Homo sapiens

<400> 237

Met Asp Leu Lys Leu Arg Ala Ala Ser Pro Ile Ile Thr Leu Val Ala
 1 5 10 15

Leu Asp Glu Ala Leu Asp Asn Tyr Thr Ile Thr Phe Leu Ile Arg Gly
 20 25 30

Val Ala Ile Gly Gln Thr Ser Leu Thr Ala Ser Val Thr Asn Lys Ala
 35 40 45

Gly Gln Arg Ile Asn Ser Ala Pro Gln Gln Ile Glu Val Phe Pro Pro
 50 55 60

Phe Arg Leu Met Pro Arg Lys Val Thr Leu Leu Ile Gly Ala Thr Met
 65 70 75 80

Gln Val Thr Ser Glu Gly Gly Pro Gln Pro Gln Ser Asn Ile Leu Phe
 85 90 95

Ser Ile Ser Asn Glu Ser Val Ala Leu Val Ser Ala Ala Gly Leu Val
 100 105 110

Gln Gly Leu Ala Ile Gly Asn Gly Thr Val Ser Gly Leu Val Gln Ala
 115 120 125

Val Asp Ala Glu Thr Gly Lys Val Val Ile Ile Ser Gln Asp Leu Val
 130 135 140

Gln Val Glu Val Leu Leu Leu Arg Ala Val Arg Ile Arg Ala Pro Ile
 145 150 155 160

Met Arg Met Arg Thr Gly Thr Gln Met Pro Ile Tyr Val Thr Gly Ile
 165 170 175

Thr Asn His Gln Asn Pro Phe Ser Phe Gly Asn Ala Val Pro Gly Leu
 180 185 190

Thr Phe His Trp Ser Val Thr Lys Arg Asp Val Leu Asp Leu Arg Gly

195	200	205
Arg His His Glu Ala Ser Ile Arg Leu Pro Ser Gln Tyr Asn Phe Ala		
210	215	220
Met Asn Val Leu Gly Arg Val Lys Gly Arg Thr Gly Leu Arg Val Val		
225	230	235 240
Val Lys Ala Val Asp Pro Thr Ser Gly Gln Leu Tyr Gly Leu Ala Arg		
	245	250 255
Glu Leu Ser Asp Glu Ile Gln Val Gln Val Phe Glu Lys Leu Gln Leu		
	260	265 270
Leu Asn Pro Glu Ile Glu Ala Glu Gln Ile Leu Met Ser Pro Asn Ser		
	275	280 285
Tyr Ile Lys Leu Gln Thr Asn Arg Asp Gly Ala Ala Ser Leu Ser Tyr		
	290	295 300
Arg Val Leu Asp Gly Pro Glu Lys Val Pro Val Val His Val Asp Glu		
305	310	315 320
Lys Gly Phe Leu Ala Ser Gly Ser Met Ile Gly Thr Ser Thr Ile Glu		
	325	330 335
Val Ile Ala Gln Glu Pro Phe Gly Ala Asn Gln Thr Ile Ile Val Ala		
	340	345 350
Val Lys Val Ser Pro Val Ser Tyr Leu Arg Val Ser Met Ser Pro Val		
	355	360 365
Leu His Thr Gln Asn Lys Glu Ala Leu Val Ala Val Pro Leu Gly Met		
	370	375 380
Thr Val Thr Phe Thr Val His Phe His Asp Asn Ser Gly Asp Val Phe		
385	390	395 400
His Ala His Ser Ser Val Leu Asn Phe Ala Thr Asn Arg Asp Asp Phe		
	405	410 415
Val Gln Ile Gly Lys Gly Pro Thr Asn Asn Thr Cys Val Val Arg Thr		
	420	425 430

Val Ser Val Gly Leu Thr Leu Leu Arg Val Trp Asp Ala Glu His Pro
 435 440 445

Gly Leu Ser Asp Phe Met Pro Leu Pro Val Leu Gln Ala Ile Ser Pro
 450 455 460

Glu Leu Ser Gly Ala Met Val Val Gly Asp Val Leu Cys Leu Ala Thr
 465 470 475 480

Val Leu Thr Ser Leu Glu Gly Leu Ser Gly Thr Trp Ser Ser Ser Ala
 485 490 495

Asn Ser Ile Leu His Ile Asp Pro Lys Thr Gly Val Ala Val Ala Arg
 500 505 510

Ala Val Gly Ser Val Thr Val Tyr Tyr Glu Val Ala Gly His Leu Arg
 515 520 525

Thr Tyr Lys Glu Val Val Val Ser Val Pro Gln Arg Ile Met Ala Arg
 530 535 540

His Leu His Pro Ile Gln Thr Ser Phe Gln Glu Ala Thr Ala Ser Lys
 545 550 555 560

Val Ile Val Ala Val Gly Asp Arg Ser Ser Asn Leu Arg Gly Glu Cys
 565 570 575

Thr Pro Thr Gln Arg Glu Val Ile Gln Ala Leu His Pro Glu Thr Leu
 580 585 590

Ile Ser Cys Gln Ser Gln Phe Lys Pro Ala Val Phe Asp Phe Pro Ser
 595 600 605

Gln Asp Val Phe Thr Val Glu Pro Gln Phe Asp Thr Ala Leu Gly Gln
 610 615 620

Tyr Phe Cys Ser Ile Thr Met His Arg Leu Thr Asp Lys Gln Arg Lys
 625 630 635 640

His Leu Ser Met Lys Lys Thr Ala Leu Val Val Ser Ala Ser Leu Ser
 645 650 655

Ser Ser His Phe Ser Thr Glu Gln Val Gly Ala Glu Val Pro Phe Ser
660 665 670

Pro Gly Leu Phe Ala Asp Gln Ala Glu Ile Leu Leu Ser Asn His Tyr
675 680 685

Thr Ser Ser Glu Ile Arg Val Phe Gly Ala Pro Glu Val Leu Glu Asn
690 695 700

Leu Glu Val Lys Ser Gly Ser Pro Ala Val Leu Ala Phe Ala Lys Glu
705 710 715 720

Lys Ser Phe Gly Trp Pro Ser Phe Ile Thr Tyr Thr Val Gly Val Leu
725 730 735

Asp Pro Ala Ala Gly Ser Gln Gly Pro Leu Ser Thr Thr Leu Thr Phe
740 745 750

Ser Ser Pro Val Thr Asn Gln Ala Ile Ala Ile Pro Val Thr Val Ala
755 760 765

Phe Val Val Asp Arg Arg Gly Pro Gly Pro Tyr Gly Ala Ser Leu Phe
770 775 780

Gln His Phe Leu Asp Ser Tyr Gln Val Met Phe Phe Thr Leu Phe Ala
785 790 795 800

Leu Leu Ala Gly Thr Ala Val Met Ile Ile Ala Tyr His Thr Val Cys
805 810 815

Thr Pro Arg Asp Leu Ala Val Pro Ala Ala Leu Thr Pro Arg Ala Ser
820 825 830

Pro Gly His Ser Pro His Tyr Phe Ala Ala Ser Ser Pro Thr Ser Pro
835 840 845

Asn Ala Leu Pro Pro Ala Arg Lys Ala Ser Pro Pro Ser Gly Leu Trp
850 855 860

Ser Pro Ala Tyr Ala Ser His
865 870

<210> 238
 <211> 728
 <212> PRT
 <213> Homo sapiens

<400> 238

Leu Pro Ala Cys Arg Leu Cys His Arg Arg Glu His Gly Arg Thr Val
 1 5 10 15

Cys Ser Gly Val Asp Thr Lys Leu Lys Phe Thr Leu Glu Pro Ser Leu
 20 25 30

Gly Gln Asn Gly Phe Gln Gln Trp Tyr Asp Ala Leu Lys Ala Val Ala
 35 40 45

Arg Leu Ser Thr Gly Ile Pro Lys Glu Trp Arg Arg Lys Val Trp Leu
 50 55 60

Thr Leu Ala Asp His Tyr Leu His Ser Ile Ala Ile Asp Trp Asp Lys
 65 70 75 80

Thr Met Arg Phe Thr Phe Asn Glu Arg Ser Asn Pro Asp Asp Asp Ser
 85 90 95

Met Gly Ile Gln Ile Val Lys Asp Leu His Arg Thr Gly Cys Ser Ser
 100 105 110

Tyr Cys Gly Gln Glu Ala Glu Gln Asp Arg Val Val Leu Lys Arg Val
 115 120 125

Leu Leu Ala Tyr Ala Arg Trp Asn Lys Thr Val Gly Tyr Cys Gln Gly
 130 135 140

Phe Asn Ile Leu Ala Ala Leu Ile Leu Glu Val Met Glu Gly Asn Glu
 145 150 155 160

Gly Asp Ala Leu Lys Ile Met Ile Tyr Leu Ile Asp Lys Val Leu Pro
 165 170 175

Glu Ser Tyr Phe Val Asn Asn Leu Arg Ala Leu Ser Val Asp Met Ala
 180 185 190

Val Phe Arg Asp Leu Leu Arg Met Lys Leu Pro Glu Leu Ser Gln His
 195 200 205

Leu Asp Thr Leu Gln Arg Thr Ala Asn Lys Glu Ser Gly Gly Gly Tyr
 210 215 220

Glu Pro Pro Leu Thr Asn Val Phe Thr Met Gln Trp Phe Leu Thr Leu
 225 230 235 240

Phe Ala Thr Cys Leu Pro Asn Gln Thr Val Leu Lys Ile Trp Asp Ser
 245 250 255

Val Phe Phe Glu Gly Ser Glu Ile Ile Leu Arg Val Ser Leu Ala Ile
 260 265 270

Trp Ala Lys Leu Gly Glu Gln Ile Glu Cys Cys Glu Thr Ala Asp Glu
 275 280 285

Phe Tyr Ser Thr Met Gly Arg Leu Thr Gln Glu Met Leu Glu Asn Asp
 290 295 300

Leu Leu Gln Ser His Glu Leu Met Gln Thr Val Tyr Ser Met Ala Pro
 305 310 315 320

Phe Pro Phe Pro Gln Leu Ala Glu Leu Arg Glu Lys Tyr Thr Tyr Asn
 325 330 335

Ile Thr Pro Phe Pro Ala Thr Val Lys Pro Thr Ser Val Ser Gly Arg
 340 345 350

His Ser Lys Ala Arg Asp Ser Asp Glu Glu Asn Asp Pro Asp Asp Glu
 355 360 365

Asp Ala Val Val Asn Ala Val Gly Cys Leu Gly Pro Phe Ser Gly Phe
 370 375 380

Leu Ala Pro Glu Leu Gln Lys Tyr Gln Lys Gln Ile Lys Glu Pro Asn
 385 390 395 400

Glu Glu Gln Ser Leu Arg Ser Asn Asn Ile Ala Glu Leu Ser Pro Gly
 405 410 415

Ala Ile Asn Ser Cys Arg Ser Glu Tyr His Ala Ala Phe Asn Ser Met
 420 425 430

Met Met Glu Arg Met Thr Thr Asp Ile Asn Ala Leu Lys Arg Gln Tyr
 435 440 445

Ser Arg Ile Lys Lys Lys Gln Gln Gln Gln Val His Gln Val Tyr Ile
 450 455 460

Arg Ala Asp Lys Gly Pro Val Thr Ser Ile Leu Pro Ser Gln Val Asn
 465 470 475 480

Ser Ser Pro Val Ile Asn His Leu Leu Leu Gly Lys Lys Met Lys Met
 485 490 495

Thr Asn Arg Ala Ala Lys Asn Ala Val Ile His Ile Pro Gly His Thr
 500 505 510

Gly Gly Lys Ile Ser Pro Val Pro Tyr Glu Asp Leu Lys Thr Lys Leu
 515 520 525

Asn Ser Pro Trp Arg Thr His Ile Arg Val His Lys Lys Asn Met Pro
 530 535 540

Arg Thr Lys Ser His Pro Gly Cys Gly Asp Thr Val Gly Leu Ile Asp
 545 550 555 560

Glu Gln Asn Glu Ala Ser Lys Thr Asn Gly Leu Gly Ala Ala Glu Ala
 565 570 575

Phe Pro Ser Gly Cys Thr Ala Thr Ala Gly Arg Glu Gly Ser Ser Pro
 580 585 590

Glu Gly Ser Thr Arg Arg Thr Ile Glu Gly Gln Ser Pro Glu Pro Val
 595 600 605

Phe Gly Asp Ala Asp Val Asp Val Ser Ala Val Gln Ala Lys Leu Gly
 610 615 620

Ala Leu Glu Leu Asn Gln Arg Asp Ala Ala Ala Glu Thr Glu Leu Arg
 625 630 635 640

Val His Pro Pro Cys Gln Arg His Cys Pro Glu Pro Pro Ser Ala Pro
 645 650 655

Glu Glu Asn Lys Ala Thr Ser Lys Ala Pro Gln Gly Ser Asn Ser Lys
 660 665 670

Thr Pro Ile Phe Ser Pro Phe Pro Ser Val Lys Pro Leu Arg Lys Ser
 675 680 685

Ala Thr Ala Arg Asn Leu Gly Leu Tyr Gly Pro Thr Glu Arg Thr Pro
 690 695 700

Thr Val His Phe Pro Gln Met Ser Arg Ser Phe Ser Lys Pro Gly Gly
 705 710 715 720

Gly Asn Ser Gly Thr Lys Lys Arg
 725

<210> 239
 <211> 787
 <212> PRT
 <213> Homo sapiens

<400> 239

Asp Leu Tyr Leu Leu Leu Leu Ser Tyr Ser Asp Lys Lys Asp His Leu
 1 5 10 15

Thr Val Glu Glu Leu Ala Gln Phe Leu Lys Val Glu Gln Lys Met Asn
 20 25 30

Asn Val Thr Thr Asp Tyr Cys Leu Asp Ile Ile Lys Lys Phe Glu Val
 35 40 45

Ser Glu Glu Asn Lys Val Lys Asn Val Leu Gly Ile Glu Gly Phe Thr
 50 55 60

Asn Phe Met Arg Ser Pro Ala Cys Asp Ile Phe Asn Pro Leu His His
 65 70 75 80

Glu Val Tyr Gln Asp Met Asp Gln Pro Leu Cys Asn Tyr Tyr Ile Ala
 85 90 95

Ser Ser His Asn Thr Tyr Leu Thr Gly Asp Gln Leu Leu Ser Gln Ser
 100 105 110

Lys Val Asp Met Tyr Ala Arg Val Leu Gln Glu Gly Cys Arg Cys Val
 115 120 125

Glu Val Asp Cys Trp Asp Gly Pro Asp Gly Glu Pro Val Val His His
 130 135 140

Gly Tyr Thr Leu Thr Ser Lys Ile Leu Phe Arg Asp Val Val Glu Thr
 145 150 155 160

Ile Asn Lys His Ala Phe Val Lys Asn Glu Phe Pro Val Ile Leu Ser
 165 170 175

Ile Glu Asn His Cys Ser Ile Gln Gln Gln Arg Lys Ile Ala Gln Tyr
 180 185 190

Leu Lys Gly Ile Phe Gly Asp Lys Leu Asp Leu Ser Ser Val Asp Thr
 195 200 205

Gly Glu Cys Lys Gln Leu Pro Ser Pro Gln Ser Leu Lys Gly Lys Ile
 210 215 220

Leu Val Lys Gly Lys Lys Leu Pro Tyr His Leu Gly Asp Asp Ala Glu
 225 230 235 240

Glu Gly Glu Val Ser Asp Glu Asp Ser Ala Asp Glu Ile Glu Asp Glu
 245 250 255

Cys Lys Phe Lys Leu His Tyr Ser Asn Gly Thr Thr Glu His Gln Val
 260 265 270

Glu Ser Phe Ile Arg Lys Lys Leu Glu Ser Leu Leu Lys Glu Ser Gln
 275 280 285

Ile Arg Asp Lys Glu Asp Pro Asp Ser Phe Thr Val Arg Ala Leu Leu
 290 295 300

Lys Ala Thr His Glu Gly Leu Asn Ala His Leu Lys Gln Ser Pro Asp
 305 310 315 320

Val Lys Glu Ser Gly Lys Lys Ser His Gly Arg Ser Leu Met Thr Asn
 325 330 335

Phe Gly Lys His Lys Lys Thr Thr Lys Ser Arg Ser Lys Ser Tyr Ser
 340 345 350

Thr Asp Asp Glu Glu Asp Thr Gln Gln Ser Thr Gly Lys Glu Gly Gly
 355 360 365

Gln Leu Tyr Arg Leu Gly Arg Arg Arg Lys Thr Met Lys Leu Cys Arg
 370 375 380

Glu Leu Ser Asp Leu Val Val Tyr Thr Asn Ser Val Ala Ala Gln Asp
 385 390 395 400

Ile Val Asp Asp Gly Thr Thr Gly Asn Val Leu Ser Phe Ser Glu Thr
 405 410 415

Arg Ala His Gln Val Val Gln Gln Lys Ser Glu Gln Phe Met Ile Tyr
 420 425 430

Asn Gln Lys Gln Leu Thr Arg Ile Tyr Pro Ser Ala Tyr Arg Ile Asp
 435 440 445

Ser Ser Asn Phe Asn Pro Leu Pro Tyr Trp Asn Ala Gly Cys Gln Leu
 450 455 460

Val Ala Leu Asn Tyr Gln Ser Glu Gly Arg Met Met Gln Leu Asn Arg
 465 470 475 480

Ala Lys Phe Lys Ala Asn Gly Asn Cys Gly Tyr Val Leu Lys Pro Gln
 485 490 495

Gln Met Cys Lys Gly Thr Phe Asn Pro Phe Ser Gly Asp Pro Leu Pro
 500 505 510

Ala Asn Pro Lys Lys Gln Leu Ile Leu Lys Val Ile Ser Gly Gln Gln
 515 520 525

Leu Pro Lys Pro Pro Asp Ser Met Phe Gly Asp Arg Gly Glu Ile Ile
 530 535 540

Asp Pro Phe Val Glu Val Glu Ile Ile Gly Leu Pro Val Asp Cys Cys
 545 550 555 560

Lys Asp Gln Thr Arg Val Val Asp Asp Asn Gly Phe Asn Pro Val Trp
 565 570 575

Glu Glu Thr Leu Thr Phe Thr Val His Met Pro Glu Ile Ala Leu Val
 580 585 590

Arg Phe Leu Val Trp Asp His Asp Pro Ile Gly Arg Asp Phe Val Gly
 595 600 605

Gln Arg Thr Val Thr Phe Ser Ser Leu Val Pro Gly Tyr Arg His Val
 610 615 620

Tyr Leu Glu Gly Leu Thr Glu Ala Ser Ile Phe Val His Ile Thr Ile
 625 630 635 640

Asn Glu Ile Tyr Gly Lys Trp Ser Pro Leu Ile Leu Asn Pro Ser Tyr
 645 650 655

Thr Ile Leu His Phe Leu Gly Ala Thr Lys Asn Arg Gln Leu Gln Gly
 660 665 670

Leu Lys Gly Leu Phe Asn Lys Asn Pro Arg His Ser Ser Ser Glu Asn
 675 680 685

Asn Ser His Tyr Val Arg Lys Arg Ser Ile Gly Asp Arg Ile Leu Arg
 690 695 700

Arg Thr Ala Ser Ala Pro Ala Lys Gly Arg Lys Lys Ser Lys Met Gly
 705 710 715 720

Phe Gln Glu Met Val Glu Ile Lys Asp Ser Val Ser Glu Ala Thr Arg
 725 730 735

Asp Gln Asp Gly Val Leu Arg Arg Thr Thr Arg Ser Leu Gln Ala Arg
 740 745 750

Pro Val Ser Met Pro Val Asp Arg Asn Leu Leu Gly Ala Leu Ser Leu
 755 760 765

Pro Val Ser Glu Thr Ala Lys Asp Ile Glu Gly Lys Glu Asn Ser Leu
 770 775 780

Val Gln Ile
 785

<210> 240

<211> 665
 <212> PRT
 <213> Homo sapiens

<400> 240

Met Ala His Glu Met Ile Gly Thr Gln Ile Val Thr Glu Arg Leu Val
 1 5 10 15

Ala Leu Leu Glu Ser Gly Thr Glu Lys Val Leu Leu Ile Asp Ser Arg
 20 25 30

Pro Phe Val Glu Tyr Asn Thr Ser His Ile Leu Glu Ala Ile Asn Ile
 35 40 45

Asn Cys Ser Lys Leu Met Lys Arg Arg Leu Gln Gln Asp Lys Val Leu
 50 55 60

Ile Thr Glu Leu Ile Gln His Ser Ala Lys His Lys Val Asp Ile Asp
 65 70 75 80

Cys Ser Gln Lys Val Val Val Tyr Asp Gln Ser Ser Gln Asp Val Ala
 85 90 95

Ser Leu Ser Ser Asp Cys Phe Leu Thr Val Leu Leu Gly Lys Leu Glu
 100 105 110

Lys Ser Phe Asn Ser Val His Leu Leu Ala Gly Gly Phe Ala Glu Phe
 115 120 125

Ser Arg Cys Phe Pro Gly Leu Cys Glu Gly Lys Ser Thr Leu Val Pro
 130 135 140

Thr Cys Ile Ser Gln Pro Cys Leu Pro Val Ala Asn Ile Gly Pro Thr
 145 150 155 160

Arg Ile Leu Pro Asn Leu Tyr Leu Gly Cys Gln Arg Asp Val Leu Asn
 165 170 175

Lys Glu Leu Met Gln Gln Asn Gly Ile Gly Tyr Val Leu Asn Ala Ser
 180 185 190

Asn Thr Cys Pro Lys Pro Asp Phe Ile Pro Glu Ser His Phe Leu Arg
 195 200 205

Val Pro Val Asn Asp Ser Phe Cys Glu Lys Ile Leu Pro Trp Leu Asp
 210 215 220

Lys Ser Val Asp Phe Ile Glu Lys Ala Lys Ala Ser Asn Gly Cys Val
 225 230 235 240

Leu Val His Cys Leu Ala Gly Ile Ser Arg Ser Ala Thr Ile Ala Ile
 245 250 255

Ala Tyr Ile Met Lys Arg Met Asp Met Ser Leu Asp Glu Ala Tyr Arg
 260 265 270

Phe Val Lys Glu Lys Arg Pro Thr Ile Ser Pro Asn Phe Asn Phe Leu
 275 280 285

Gly Gln Leu Leu Asp Tyr Glu Lys Lys Ile Lys Asn Gln Thr Gly Ala
 290 295 300

Ser Gly Pro Lys Ser Lys Leu Lys Leu Leu His Leu Glu Lys Pro Asn
 305 310 315 320

Glu Pro Val Pro Ala Val Ser Glu Gly Gly Gln Lys Ser Glu Thr Pro
 325 330 335

Leu Ser Pro Pro Cys Ala Asp Ser Ala Thr Ser Glu Ala Ala Gly Gln
 340 345 350

Arg Pro Val His Pro Ala Ser Val Pro Ser Val Pro Ser Val Gln Pro
 355 360 365

Ser Leu Leu Glu Asp Ser Pro Leu Val Gln Ala Leu Ser Gly Leu His
 370 375 380

Leu Ser Ala Asp Arg Leu Glu Asp Ser Asn Lys Leu Lys Arg Ser Phe
 385 390 395 400

Ser Leu Asp Ile Lys Ser Val Ser Tyr Ser Ala Ser Met Ala Ala Ser
 405 410 415

Leu His Gly Phe Ser Ser Ser Glu Asp Ala Leu Glu Tyr Tyr Lys Pro
 420 425 430

Ser Thr Thr Leu Asp Gly Thr Asn Lys Leu Cys Gln Phe Ser Pro Val
 435 440 445

Gln Glu Leu Ser Glu Gln Thr Pro Glu Thr Ser Pro Asp Lys Glu Glu
 450 455 460

Ala Ser Ile Pro Lys Lys Leu Gln Thr Ala Arg Pro Ser Asp Ser Gln
 465 470 475 480

Ser Lys Arg Leu His Ser Val Arg Thr Ser Ser Ser Gly Thr Ala Gln
 485 490 495

Arg Ser Leu Leu Ser Pro Leu His Arg Ser Gly Ser Val Glu Asp Asn
 500 505 510

Tyr His Thr Ser Phe Leu Phe Gly Leu Ser Thr Ser Gln Gln His Leu
 515 520 525

Thr Lys Ser Ala Gly Leu Gly Leu Lys Gly Trp His Ser Asp Ile Leu
 530 535 540

Ala Pro Gln Thr Ser Thr Pro Ser Leu Thr Ser Ser Trp Tyr Phe Ala
 545 550 555 560

Thr Glu Ser Ser His Phe Tyr Ser Ala Ser Ala Ile Tyr Gly Gly Ser
 565 570 575

Ala Ser Tyr Ser Ala Tyr Ser Cys Ser Gln Leu Pro Thr Cys Gly Asp
 580 585 590

Gln Val Tyr Ser Val Arg Arg Arg Gln Lys Pro Ser Asp Arg Ala Asp
 595 600 605

Ser Arg Arg Ser Trp His Glu Glu Ser Pro Phe Glu Lys Gln Phe Lys
 610 615 620

Arg Arg Ser Cys Gln Met Glu Phe Gly Glu Ser Ile Met Ser Glu Asn
 625 630 635 640

Arg Ser Arg Glu Glu Leu Gly Lys Val Gly Ser Gln Ser Ser Phe Ser
 645 650 655

Gly Ser Met Glu Ile Ile Glu Val Ser

660

665

<210> 241
 <211> 563
 <212> PRT
 <213> Homo sapiens

<400> 241

Met Trp Ala Val Leu Arg Leu Ala Leu Arg Pro Cys Ala Arg Ala Ser
 1 5 10 15

Pro Ala Gly Pro Arg Ala Tyr His Gly Asp Ser Val Ala Ser Leu Gly
 20 25 30

Thr Gln Pro Asp Leu Gly Ser Ala Leu Tyr Gln Glu Asn Tyr Lys Gln
 35 40 45

Met Lys Ala Leu Val Asn Gln Leu His Glu Arg Val Glu His Ile Lys
 50 55 60

Leu Gly Gly Gly Glu Lys Ala Arg Ala Leu His Ile Ser Arg Gly Lys
 65 70 75 80

Leu Leu Pro Arg Glu Arg Ile Asp Asn Leu Ile Asp Pro Gly Ser Pro
 85 90 95

Phe Leu Glu Leu Ser Gln Phe Ala Gly Tyr Gln Leu Tyr Asp Asn Glu
 100 105 110

Glu Val Pro Gly Gly Gly Ile Ile Thr Gly Ile Gly Arg Val Ser Gly
 115 120 125

Val Glu Cys Met Ile Ile Ala Asn Asp Ala Thr Val Lys Gly Gly Ala
 130 135 140

Tyr Tyr Pro Val Thr Val Lys Lys Gln Leu Arg Ala Gln Glu Ile Ala
 145 150 155 160

Met Gln Asn Arg Leu Pro Cys Ile Tyr Leu Val Asp Ser Gly Gly Ala
 165 170 175

Tyr Leu Pro Arg Gln Ala Asp Val Phe Pro Asp Arg Asp His Phe Gly
 180 185 190

Arg Thr Phe Tyr Asn Gln Ala Ile Met Ser Ser Lys Asn Ile Ala Gln
 195 200 205

Ile Ala Val Val Met Gly Ser Cys Thr Ala Gly Gly Ala Tyr Val Pro
 210 215 220

Ala Met Ala Asp Glu Asn Ile Ile Val Arg Lys Gln Gly Thr Ile Phe
 225 230 235 240

Leu Ala Gly Pro Pro Leu Val Lys Ala Ala Thr Gly Glu Glu Val Ser
 245 250 255

Ala Glu Asp Leu Gly Gly Ala Asp Leu His Cys Arg Lys Ser Gly Val
 260 265 270

Ser Asp His Trp Ala Leu Asp Asp His His Ala Leu His Leu Thr Arg
 275 280 285

Lys Val Val Arg Asn Leu Asn Tyr Gln Lys Lys Leu Asp Val Thr Ile
 290 295 300

Glu Pro Ser Glu Glu Pro Leu Phe Pro Ala Asp Glu Leu Tyr Gly Ile
 305 310 315 320

Val Gly Ala Asn Leu Lys Arg Ser Phe Asp Val Arg Glu Val Ile Ala
 325 330 335

Arg Ile Val Asp Gly Ser Arg Phe Thr Glu Phe Lys Ala Phe Tyr Gly
 340 345 350

Asp Thr Leu Val Thr Gly Phe Ala Arg Ile Phe Gly Tyr Pro Val Gly
 355 360 365

Ile Val Gly Asn Asn Gly Val Leu Phe Ser Glu Ser Ala Lys Lys Gly
 370 375 380

Thr His Phe Val Gln Leu Cys Cys Gln Arg Asn Ile Pro Leu Leu Phe
 385 390 395 400

Leu Gln Asn Ile Thr Gly Phe Met Val Gly Arg Glu Tyr Glu Ala Glu
 405 410 415

Gly Ile Ala Lys Asp Gly Ala Lys Met Val Ala Ala Val Ala Cys Ala
 420 425 430

Gln Val Pro Lys Ile Thr Leu Ile Ile Gly Gly Ser Tyr Gly Ala Gly
 435 440 445

Asn Tyr Gly Met Cys Gly Arg Ala Tyr Ser Pro Arg Phe Leu Tyr Ile
 450 455 460

Trp Pro Asn Ala Arg Ile Ser Val Met Gly Gly Glu Gln Ala Ala Asn
 465 470 475 480

Val Leu Ala Thr Ile Thr Lys Asp Gln Arg Ala Arg Glu Gly Lys Gln
 485 490 495

Phe Ser Ser Ala Asp Glu Ala Ala Leu Lys Glu Pro Ile Ile Lys Lys
 500 505 510

Phe Glu Glu Glu Gly Asn Pro Tyr Tyr Ser Ser Ala Arg Val Trp Asp
 515 520 525

Asp Gly Ile Ile Asp Pro Ala Asp Thr Arg Leu Val Leu Gly Leu Ser
 530 535 540

Phe Ser Ala Ala Leu Asn Ala Pro Ile Glu Lys Thr Asp Phe Gly Ile
 545 550 555 560

Phe Arg Met

<210> 242
 <211> 758
 <212> PRT
 <213> Homo sapiens

<400> 242

Met Ala Glu Pro Arg Gln Glu Phe Glu Val Met Glu Asp His Ala Gly
 1 5 10 15

Thr Tyr Gly Leu Gly Asp Arg Lys Asp Gln Gly Gly Tyr Thr Met His
 20 25 30

Gln Asp Gln Glu Gly Asp Thr Asp Ala Gly Leu Lys Glu Ser Pro Leu
 35 40 45

Gln Thr Pro Thr Glu Asp Gly Ser Glu Glu Pro Gly Ser Glu Thr Ser
50 55 60

Asp Ala Lys Ser Thr Pro Thr Ala Glu Asp Val Thr Ala Pro Leu Val
65 70 75 80

Asp Glu Gly Ala Pro Gly Lys Gln Ala Ala Ala Gln Pro His Thr Glu
85 90 95

Ile Pro Glu Gly Thr Thr Ala Glu Glu Ala Gly Ile Gly Asp Thr Pro
100 105 110

Ser Leu Glu Asp Glu Ala Ala Gly His Val Thr Gln Glu Pro Glu Ser
115 120 125

Gly Lys Val Val Gln Glu Gly Phe Leu Arg Glu Pro Gly Pro Pro Gly
130 135 140

Leu Ser His Gln Leu Met Ser Gly Met Pro Gly Ala Pro Leu Leu Pro
145 150 155 160

Glu Gly Pro Arg Glu Ala Thr Arg Gln Pro Ser Gly Thr Gly Pro Glu
165 170 175

Asp Thr Glu Gly Gly Arg His Ala Pro Glu Leu Leu Lys His Gln Leu
180 185 190

Leu Gly Asp Leu His Gln Glu Gly Pro Pro Leu Lys Gly Ala Gly Gly
195 200 205

Lys Glu Arg Pro Gly Ser Lys Glu Glu Val Asp Glu Asp Arg Asp Val
210 215 220

Asp Glu Ser Ser Pro Gln Asp Ser Pro Pro Ser Lys Ala Ser Pro Ala
225 230 235 240

Gln Asp Gly Arg Pro Pro Gln Thr Ala Ala Arg Glu Ala Thr Ser Ile
245 250 255

Pro Gly Phe Pro Ala Glu Gly Ala Ile Pro Leu Pro Val Asp Phe Leu
260 265 270

Ser Lys Val Ser Thr Glu Ile Pro Ala Ser Glu Pro Asp Gly Pro Ser
 275 280 285

Val Gly Arg Ala Lys Gly Gln Asp Ala Pro Leu Glu Phe Thr Phe His
 290 295 300

Val Glu Ile Thr Pro Asn Val Gln Lys Glu Gln Ala His Ser Glu Glu
 305 310 315 320

His Leu Gly Arg Ala Ala Phe Pro Gly Ala Pro Gly Glu Gly Pro Glu
 325 330 335

Ala Arg Gly Pro Ser Leu Gly Glu Asp Thr Lys Glu Ala Asp Leu Pro
 340 345 350

Glu Pro Ser Glu Lys Gln Pro Ala Ala Ala Pro Arg Gly Lys Pro Val
 355 360 365

Ser Arg Val Pro Gln Leu Lys Ala Arg Met Val Ser Lys Ser Lys Asp
 370 375 380

Gly Thr Gly Ser Asp Asp Lys Lys Ala Lys Thr Ser Thr Arg Ser Ser
 385 390 395 400

Ala Lys Thr Leu Lys Asn Arg Pro Cys Leu Ser Pro Lys Leu Pro Thr
 405 410 415

Pro Gly Ser Ser Asp Pro Leu Ile Gln Pro Ser Ser Pro Ala Val Cys
 420 425 430

Pro Glu Pro Pro Ser Ser Pro Lys His Val Ser Ser Val Thr Ser Arg
 435 440 445

Thr Gly Ser Ser Gly Ala Lys Glu Met Lys Leu Lys Gly Ala Asp Gly
 450 455 460

Lys Thr Lys Ile Ala Thr Pro Arg Gly Ala Ala Pro Pro Gly Gln Lys
 465 470 475 480

Gly Gln Ala Asn Ala Thr Arg Ile Pro Ala Lys Thr Pro Pro Ala Pro
 485 490 495

Lys Thr Pro Pro Ser Ser Gly Glu Pro Pro Lys Ser Gly Asp Arg Ser
500 505 510

Gly Tyr Ser Ser Pro Gly Ser Pro Gly Thr Pro Gly Ser Arg Ser Arg
515 520 525

Thr Pro Ser Leu Pro Thr Pro Pro Thr Arg Glu Pro Lys Lys Val Ala
530 535 540

Val Val Arg Thr Pro Pro Lys Ser Pro Ser Ser Ala Lys Ser Arg Leu
545 550 555 560

Gln Thr Ala Pro Val Pro Met Pro Asp Leu Lys Asn Val Lys Ser Lys
565 570 575

Ile Gly Ser Thr Glu Asn Leu Lys His Gln Pro Gly Gly Gly Lys Val
580 585 590

Gln Ile Ile Asn Lys Lys Leu Asp Leu Ser Asn Val Gln Ser Lys Cys
595 600 605

Gly Ser Lys Asp Asn Ile Lys His Val Pro Gly Gly Gly Ser Val Gln
610 615 620

Ile Val Tyr Lys Pro Val Asp Leu Ser Lys Val Thr Ser Lys Cys Gly
625 630 635 640

Ser Leu Gly Asn Ile His His Lys Pro Gly Gly Gly Gln Val Glu Val
645 650 655

Lys Ser Glu Lys Leu Asp Phe Lys Asp Arg Val Gln Ser Lys Ile Gly
660 665 670

Ser Leu Asp Asn Ile Thr His Val Pro Gly Gly Gly Asn Lys Lys Ile
675 680 685

Glu Thr His Lys Leu Thr Phe Arg Glu Asn Ala Lys Ala Lys Thr Asp
690 695 700

His Gly Ala Glu Ile Val Tyr Lys Ser Pro Val Val Ser Gly Asp Thr
705 710 715 720

Ser Pro Arg His Leu Ser Asn Val Ser Ser Thr Gly Ser Ile Asp Met

725

730

735

Val Asp Ser Pro Gln Leu Ala Thr Leu Ala Asp Glu Val Ser Ala Ser
 740 745 750

Leu Ala Lys Gln Gly Leu
 755

<210> 243
 <211> 547
 <212> PRT
 <213> Homo sapiens

<400> 243

Met Glu Asn Asp Glu Ser Ala Lys Glu Glu Lys Ser Asp Leu Lys Glu
 1 5 10 15

Lys Ser Thr Gly Ser Lys Lys Ala Asn Arg Phe His Pro Tyr Ser Lys
 20 25 30

Asp Lys Asn Ser Gly Thr Gly Glu Lys Lys Gly Pro Asn Arg Asn Arg
 35 40 45

Val Phe Ile Ser Asn Ile Pro Tyr Asp Met Lys Trp Gln Ala Ile Lys
 50 55 60

Asp Leu Met Arg Glu Lys Val Gly Glu Val Thr Tyr Val Glu Leu Phe
 65 70 75 80

Lys Asp Ala Glu Gly Lys Ser Arg Gly Cys Gly Val Val Glu Phe Lys
 85 90 95

Asp Glu Glu Phe Val Lys Lys Ala Leu Glu Thr Met Asn Lys Tyr Asp
 100 105 110

Leu Ser Gly Arg Arg Val Asn Ile Lys Glu Asp Pro Asp Gly Glu Asn
 115 120 125

Ala Arg Arg Ala Leu Gln Arg Thr Gly Thr Ser Phe Gln Gly Ser His
 130 135 140

Ala Ser Asp Val Gly Ser Gly Leu Val Asn Leu Pro Pro Ser Ile Leu
 145 150 155 160

Asn Asn Pro Asn Ile Pro Pro Glu Val Ile Ser Asn Leu Gln Ala Gly
 165 170 175

Arg Leu Gly Ser Thr Ile Phe Val Ala Asn Leu Asp Phe Lys Val Gly
 180 185 190

Trp Lys Lys Leu Lys Glu Val Phe Ser Ile Ala Gly Thr Val Lys Ala
 195 200 205

Gly Ser Tyr Lys Glu Asp Lys Asp Gly Lys Ser Arg Gly Met Gly Thr
 210 215 220

Val Thr Phe Glu Gln Ala Ile Glu Ala Val Gln Ala Ile Ser Met Phe
 225 230 235 240

Asn Gly Gln Phe Leu Phe Asp Arg Pro Met His Val Lys Met Asp Asp
 245 250 255

Lys Ser Val Pro His Glu Glu Tyr Arg Ser Pro Asp Gly Lys Thr Pro
 260 265 270

Gln Leu Pro Arg Gly Leu Gly Gly Ile Gly Met Gly Leu Gly Pro Gly
 275 280 285

Gly Gln Pro Ile Ser Ala Ser Gln Leu Asn Ile Gly Gly Val Met Gly
 290 295 300

Asn Leu Gly Pro Gly Gly Met Gly Met Asp Gly Pro Gly Phe Gly Gly
 305 310 315 320

Met Asn Arg Ile Gly Gly Gly Ile Gly Phe Gly Gly Leu Glu Ala Met
 325 330 335

Asn Ser Met Gly Gly Phe Gly Gly Val Gly Arg Met Gly Glu Leu Tyr
 340 345 350

Arg Gly Ala Met Thr Ser Ser Met Glu Arg Asp Phe Gly His Arg Asp
 355 360 365

Ile Gly Leu Ser Arg Gly Phe Gly Asp Ser Phe Gly Arg Leu Gly Ser
 370 375 380

Ala Met Ile Gly Gly Ile Thr Gly Arg Ile Gly Ser Ser Asn Met Gly
385 390 395 400

Pro Val Gly Ser Gly Ile Ser Gly Gly Met Gly Ser Met Asn Ser Val
405 410 415

Thr Gly Gly Met Gly Met Gly Leu Asp Arg Met Ser Ser Ser Phe Asp
420 425 430

Arg Met Gly Pro Gly Ile Gly Ala Ile Leu Glu Arg Ser Ile Asp Met
435 440 445

Asp Arg Gly Phe Leu Ser Gly Pro Met Gly Ser Gly Met Arg Glu Arg
450 455 460

Ile Gly Ser Lys Gly Asn Gln Ile Phe Val Arg Asn Leu Pro Phe Asp
465 470 475 480

Leu Thr Trp Gln Lys Leu Lys Glu Lys Phe Ser Gln Cys Gly His Val
485 490 495

Met Phe Ala Glu Ile Lys Met Glu Asn Gly Lys Ser Lys Gly Cys Gly
500 505 510

Thr Val Arg Phe Asp Ser Pro Glu Ser Ala Glu Lys Ala Cys Arg Ile
515 520 525

Met Asn Gly Ile Lys Ile Ser Gly Arg Glu Ile Asp Val Arg Leu Asp
530 535 540

Arg Asn Ala
545

<210> 244
<211> 1022
<212> PRT
<213> Homo sapiens

<400> 244

Met Asn Asn Asn Trp Asn Val Cys Phe Phe Leu Phe Cys Pro Ser Ile
1 5 10 15

Thr Arg Thr Phe Ala Ser Gly Lys Thr Glu Lys Val Ile Phe Gln Ala
20 25 30

Leu Lys Glu Leu Gly Leu Pro Ser Gly Lys Asn Asp Glu Ile Glu Pro
 35 40 45

Thr Ala Phe Ser Tyr Glu Lys Phe Tyr Glu Leu Thr Gln Lys Ile Cys
 50 55 60

Pro Arg Thr Asp Ile Glu Asp Leu Phe Lys Lys Ile Asn Gly Asp Lys
 65 70 75 80

Thr Asp Tyr Leu Thr Val Asp Gln Leu Val Ser Phe Leu Asn Glu His
 85 90 95

Gln Arg Asp Pro Arg Leu Asn Glu Ile Leu Phe Pro Phe Tyr Asp Ala
 100 105 110

Lys Arg Ala Met Gln Ile Ile Glu Met Tyr Glu Pro Asp Glu Asp Leu
 115 120 125

Lys Lys Lys Gly Leu Ile Ser Ser Asp Gly Phe Cys Arg Tyr Leu Met
 130 135 140

Ser Asp Glu Asn Ala Pro Val Phe Leu Asp Arg Leu Glu Leu Tyr Gln
 145 150 155 160

Glu Met Asp His Pro Leu Ala His Tyr Phe Ile Ser Ser Ser His Asn
 165 170 175

Thr Tyr Leu Thr Gly Arg Gln Phe Gly Gly Lys Ser Ser Val Glu Met
 180 185 190

Tyr Arg Gln Val Leu Leu Ala Gly Cys Arg Cys Val Glu Leu Asp Cys
 195 200 205

Trp Asp Gly Lys Gly Glu Asp Gln Glu Pro Ile Ile Thr His Gly Lys
 210 215 220

Ala Met Cys Thr Asp Ile Leu Phe Lys Asp Val Ile Gln Ala Ile Lys
 225 230 235 240

Glu Thr Ala Phe Val Thr Ser Glu Tyr Pro Val Ile Leu Ser Phe Glu
 245 250 255

Asn His Cys Ser Lys Tyr Gln Gln Tyr Lys Met Ser Lys Tyr Cys Glu
 260 265 270

Asp Leu Phe Gly Asp Leu Leu Leu Lys Gln Ala Leu Glu Ser His Pro
 275 280 285

Leu Glu Pro Gly Arg Pro Leu Pro Ser Pro Asn Asp Leu Lys Arg Lys
 290 295 300

Ile Leu Ile Lys Asn Lys Arg Leu Lys Pro Glu Val Glu Lys Lys Gln
 305 310 315 320

Leu Glu Ala Leu Arg Ser Met Met Glu Ala Gly Glu Ser Ala Ser Pro
 325 330 335

Ala Asn Ile Leu Glu Asp Asp Asn Glu Glu Glu Ile Glu Ser Ala Asp
 340 345 350

Gln Glu Glu Glu Ala His Pro Glu Phe Lys Phe Gly Asn Glu Leu Ser
 355 360 365

Ala Asp Asp Leu Gly His Lys Glu Ala Val Ala Asn Ser Val Lys Lys
 370 375 380

Gly Leu Val Thr Val Glu Asp Glu Gln Ala Trp Met Ala Ser Tyr Lys
 385 390 395 400

Tyr Val Gly Ala Thr Thr Asn Ile His Pro Tyr Leu Ser Thr Met Ile
 405 410 415

Asn Tyr Ala Gln Pro Val Lys Phe Gln Gly Phe His Val Ala Glu Glu
 420 425 430

Arg Asn Ile His Tyr Asn Met Ser Ser Phe Asn Glu Ser Val Gly Leu
 435 440 445

Gly Tyr Leu Lys Thr His Ala Ile Glu Phe Val Asn Tyr Asn Lys Arg
 450 455 460

Gln Met Ser Arg Ile Tyr Pro Lys Gly Gly Arg Val Asp Ser Ser Asn
 465 470 475 480

Tyr Met Pro Gln Ile Phe Trp Asn Ala Gly Cys Gln Met Val Ser Leu
 485 490 495

Asn Tyr Gln Thr Pro Asp Leu Ala Met Gln Leu Asn Gln Gly Lys Phe
 500 505 510

Glu Tyr Asn Gly Ser Cys Gly Tyr Leu Leu Lys Pro Asp Phe Met Arg
 515 520 525

Arg Pro Asp Arg Thr Phe Asp Pro Phe Ser Glu Thr Pro Val Asp Gly
 530 535 540

Val Ile Ala Ala Thr Cys Ser Val Gln Val Ile Ser Gly Gln Phe Leu
 545 550 555 560

Ser Asp Lys Lys Ile Gly Thr Tyr Val Glu Val Asp Met Tyr Gly Leu
 565 570 575

Pro Thr Asp Thr Ile Arg Lys Glu Phe Arg Thr Arg Met Val Met Asn
 580 585 590

Asn Gly Leu Asn Pro Val Tyr Asn Glu Glu Ser Leu Val Phe Arg Lys
 595 600 605

Val Ile Leu Pro Asp Leu Ala Val Leu Arg Ile Ala Val Tyr Asp Asp
 610 615 620

Asn Asn Lys Leu Ile Gly Gln Arg Ile Pro Pro Leu Asp Gly Leu Gln
 625 630 635 640

Ala Gly Tyr Arg His Ile Ser Leu Arg Asn Glu Gly Asn Lys Pro Leu
 645 650 655

Ser Leu Pro Thr Ile Phe Cys Asn Ile Val Leu Lys Thr Tyr Val Pro
 660 665 670

Asp Gly Phe Gly Asp Ile Val Asp Ala Leu Ser Asp Pro Lys Thr Phe
 675 680 685

Leu Ser Ile Thr Glu Lys Arg Ala Asp Gln Met Arg Ala Met Gly Ile
 690 695 700

Glu Thr Ser Asp Ile Ala Asp Val Pro Ser Asp Thr Ser Lys Asn Asp

705	710	715	720
Lys Lys Gly Lys Ala Asn Thr Ala Lys Ala Asn Val Thr Pro Gln Ser	725	730	735
Ser Ser Glu Leu Arg Pro Thr Thr Thr Ala Ala Leu Pro Ser Gly Val	740	745	750
Glu Ala Lys Lys Gly Ile Glu Leu Ile Pro Gln Val Arg Ile Glu Asp	755	760	765
Leu Lys Gln Met Lys Ala Tyr Leu Lys His Leu Lys Lys Gln Gln Lys	770	775	780
Glu Leu Asn Ser Leu Lys Lys Lys His Ala Lys Glu His Ser Thr Met	785	790	795
Gln Lys Leu His Cys Thr Gln Val Asp Lys Ile Val Ala Gln Tyr Asp	805	810	815
Lys Glu Lys Ser Thr His Glu Lys Ile Leu Glu Lys Ala Met Lys Lys	820	825	830
Lys Gly Gly Ser Asn Cys Leu Glu Met Lys Lys Glu Thr Glu Ile Lys	835	840	845
Ile Gln Thr Leu Thr Ser Asp His Lys Ser Lys Val Lys Glu Ile Val	850	855	860
Ala Gln His Thr Lys Glu Trp Ser Glu Met Ile Asn Thr His Ser Ala	865	870	875
Glu Glu Gln Glu Ile Arg Asp Leu His Leu Ser Gln Gln Cys Glu Leu	885	890	895
Leu Lys Lys Leu Leu Ile Asn Ala His Glu Gln Gln Thr Gln Gln Leu	900	905	910
Lys Leu Ser His Asp Arg Glu Ser Lys Glu Met Arg Ala His Gln Ala	915	920	925
Lys Ile Ser Met Glu Asn Ser Lys Ala Ile Ser Gln Asp Lys Ser Ile	930	935	940

Lys Asn Lys Ala Glu Arg Glu Arg Arg Val Arg Glu Leu Asn Ser Ser
 945 950 955 960

Asn Thr Lys Lys Phe Leu Glu Glu Arg Lys Arg Leu Ala Met Lys Gln
 965 970 975

Ser Lys Glu Met Asp Gln Leu Lys Lys Val Gln Leu Glu His Leu Glu
 980 985 990

Phe Leu Glu Lys Gln Asn Glu Gln Ala Lys Glu Met Gln Gln Met Val
 995 1000 1005

Lys Leu Glu Ala Glu Met Asp Arg Arg Pro Ala Thr Val Val
 1010 1015 1020

<210> 245
 <211> 335
 <212> PRT
 <213> Homo sapiens

<400> 245

Met Gly Ser Ala Ser Pro Gly Leu Ser Ser Val Ser Pro Ser His Leu
 1 5 10 15

Leu Leu Pro Pro Asp Thr Val Ser Arg Thr Gly Leu Glu Lys Ala Ala
 20 25 30

Ala Gly Ala Val Gly Leu Glu Arg Arg Asp Trp Ser Pro Ser Pro Pro
 35 40 45

Ala Thr Pro Glu Gln Gly Leu Ser Ala Phe Tyr Leu Ser Tyr Phe Asp
 50 55 60

Met Leu Tyr Pro Glu Asp Ser Ser Trp Ala Ala Lys Ala Pro Gly Ala
 65 70 75 80

Ser Ser Arg Glu Glu Pro Pro Glu Glu Pro Glu Gln Cys Pro Val Ile
 85 90 95

Asp Ser Gln Ala Pro Ala Gly Ser Leu Asp Leu Val Pro Gly Gly Leu
 100 105 110

Thr Leu Glu Glu His Ser Leu Glu Gln Val Gln Ser Met Val Val Gly
 115 120 125

Glu Val Leu Lys Asp Ile Glu Thr Ala Cys Lys Leu Leu Asn Ile Thr
 130 135 140

Ala Asp Pro Met Asp Trp Ser Pro Ser Asn Val Gln Lys Trp Leu Leu
 145 150 155 160

Trp Thr Glu His Gln Tyr Arg Leu Pro Pro Met Gly Lys Ala Phe Gln
 165 170 175

Glu Leu Ala Gly Lys Glu Leu Cys Ala Met Ser Glu Glu Gln Phe Arg
 180 185 190

Gln Arg Ser Pro Leu Gly Gly Asp Val Leu His Ala His Leu Asp Ile
 195 200 205

Trp Lys Ser Ala Ala Trp Met Lys Glu Arg Thr Ser Pro Gly Ala Ile
 210 215 220

His Tyr Cys Ala Ser Thr Ser Glu Glu Ser Trp Thr Asp Ser Glu Val
 225 230 235 240

Asp Ser Ser Cys Ser Gly Gln Pro Ile His Leu Trp Gln Phe Leu Lys
 245 250 255

Glu Leu Leu Leu Lys Pro His Ser Tyr Gly Arg Phe Ile Arg Trp Leu
 260 265 270

Asn Lys Glu Lys Gly Ile Phe Lys Ile Glu Asp Ser Ala Gln Val Ala
 275 280 285

Arg Leu Trp Gly Ile Arg Lys Asn Arg Pro Ala Met Asn Tyr Asp Lys
 290 295 300

Leu Ser Arg Ser Ile Arg Gln Tyr Tyr Lys Lys Gly Ile Ile Arg Lys
 305 310 315 320

Pro Asp Ile Ser Gln Arg Leu Val Tyr Gln Phe Val His Pro Ile
 325 330 335

<210> 246

<211> 174
 <212> PRT
 <213> Homo sapiens

<400> 246

Met Ala Ala Ala Met Val Pro Gly Arg Ser Glu Ser Trp Glu Arg Gly
 1 5 10 15

Glu Pro Gly Arg Pro Ala Leu Tyr Phe Cys Gly Ser Ile Arg Gly Gly
 20 25 30

Arg Glu Asp Arg Thr Leu Tyr Glu Arg Ile Val Ser Arg Leu Arg Arg
 35 40 45

Phe Gly Thr Val Leu Thr Glu His Val Ala Ala Ala Glu Leu Gly Ala
 50 55 60

Arg Gly Glu Glu Ala Ala Gly Gly Asp Arg Leu Ile His Glu Gln Asp
 65 70 75 80

Leu Glu Trp Leu Gln Gln Ala Asp Val Val Val Ala Glu Val Thr Gln
 85 90 95

Pro Ser Leu Gly Val Gly Tyr Glu Leu Gly Arg Ala Val Ala Phe Asn
 100 105 110

Lys Arg Ile Leu Cys Leu Phe Arg Pro Gln Ser Gly Arg Val Leu Ser
 115 120 125

Ala Met Ile Arg Gly Ala Ala Asp Gly Ser Arg Phe Gln Val Trp Asp
 130 135 140

Tyr Glu Glu Gly Glu Val Glu Ala Leu Leu Asp Arg Tyr Phe Glu Ala
 145 150 155 160

Asp Pro Pro Gly Gln Val Ala Ala Ser Pro Asp Pro Thr Thr
 165 170

<210> 247
 <211> 665
 <212> PRT
 <213> Homo sapiens

<400> 247

Met Ala His Glu Met Ile Gly Thr Gln Ile Val Thr Glu Arg Leu Val
1 5 10 15

Ala Leu Leu Glu Ser Gly Thr Glu Lys Val Leu Leu Ile Asp Ser Arg
20 25 30

Pro Phe Val Glu Tyr Asn Thr Ser His Ile Leu Glu Ala Ile Asn Ile
35 40 45

Asn Cys Ser Lys Leu Met Lys Arg Arg Leu Gln Gln Asp Lys Val Leu
50 55 60

Ile Thr Glu Leu Ile Gln His Ser Ala Lys His Lys Val Asp Ile Asp
65 70 75 80

Cys Ser Gln Lys Val Val Val Tyr Asp Gln Ser Ser Gln Asp Val Ala
85 90 95

Ser Leu Ser Ser Asp Cys Phe Leu Thr Val Leu Leu Gly Lys Leu Glu
100 105 110

Lys Ser Phe Asn Ser Val His Leu Leu Ala Gly Gly Phe Ala Glu Phe
115 120 125

Ser Arg Cys Phe Pro Gly Leu Cys Glu Gly Lys Ser Thr Leu Val Pro
130 135 140

Thr Cys Ile Ser Gln Pro Cys Leu Pro Val Ala Asn Ile Gly Pro Thr
145 150 155 160

Arg Ile Leu Pro Asn Leu Tyr Leu Gly Cys Gln Arg Asp Val Leu Asn
165 170 175

Lys Glu Leu Met Gln Gln Asn Gly Ile Gly Tyr Val Leu Asn Ala Ser
180 185 190

Asn Thr Cys Pro Lys Pro Asp Phe Ile Pro Glu Ser His Phe Leu Arg
195 200 205

Val Pro Val Asn Asp Ser Phe Cys Glu Lys Ile Leu Pro Trp Leu Asp
210 215 220

Lys Ser Val Asp Phe Ile Glu Lys Ala Lys Ala Ser Asn Gly Cys Val

225 230 235 240
 Leu Val His Cys Leu Ala Gly Ile Ser Arg Ser Ala Thr Ile Ala Ile
 245 250 255
 Ala Tyr Ile Met Lys Arg Met Asp Met Ser Leu Asp Glu Ala Tyr Arg
 260 265 270
 Phe Val Lys Glu Lys Arg Pro Thr Ile Ser Pro Asn Phe Asn Phe Leu
 275 280 285
 Gly Gln Leu Leu Asp Tyr Glu Lys Lys Ile Lys Asn Gln Thr Gly Ala
 290 295 300
 Ser Gly Pro Lys Ser Lys Leu Lys Leu Leu His Leu Glu Lys Pro Asn
 305 310 315 320
 Glu Pro Val Pro Ala Val Ser Glu Gly Gly Gln Lys Ser Glu Thr Pro
 325 330 335
 Leu Ser Pro Pro Cys Ala Asp Ser Ala Thr Ser Glu Ala Ala Gly Gln
 340 345 350
 Arg Pro Val His Pro Ala Ser Val Pro Ser Val Pro Ser Val Gln Pro
 355 360 365
 Ser Leu Leu Glu Asp Ser Pro Leu Val Gln Ala Leu Ser Gly Leu His
 370 375 380
 Leu Ser Ala Asp Arg Leu Glu Asp Ser Asn Lys Leu Lys Arg Ser Phe
 385 390 395 400
 Ser Leu Asp Ile Lys Ser Val Ser Tyr Ser Ala Ser Met Ala Ala Ser
 405 410 415
 Leu His Gly Phe Ser Ser Ser Glu Asp Ala Leu Glu Tyr Tyr Lys Pro
 420 425 430
 Ser Thr Thr Leu Asp Gly Thr Asn Lys Leu Cys Gln Phe Ser Pro Val
 435 440 445
 Gln Glu Leu Ser Glu Gln Thr Pro Glu Thr Ser Pro Asp Lys Glu Glu
 450 455 460

Ala Ser Ile Pro Lys Lys Leu Gln Thr Ala Arg Pro Ser Asp Ser Gln
465 470 475 480

Ser Lys Arg Leu His Ser Val Arg Thr Ser Ser Ser Gly Thr Ala Gln
485 490 495

Arg Ser Leu Leu Ser Pro Leu His Arg Ser Gly Ser Val Glu Asp Asn
500 505 510

Tyr His Thr Ser Phe Leu Phe Gly Leu Ser Thr Ser Gln Gln His Leu
515 520 525

Thr Lys Ser Ala Gly Leu Gly Leu Lys Gly Trp His Ser Asp Ile Leu
530 535 540

Ala Pro Gln Thr Ser Thr Pro Ser Leu Thr Ser Ser Trp Tyr Phe Ala
545 550 555 560

Thr Glu Ser Ser His Phe Tyr Ser Ala Ser Ala Ile Tyr Gly Gly Ser
565 570 575

Ala Ser Tyr Ser Ala Tyr Ser Cys Ser Gln Leu Pro Thr Cys Gly Asp
580 585 590

Gln Val Tyr Ser Val Arg Arg Arg Gln Lys Pro Ser Asp Arg Ala Asp
595 600 605

Ser Arg Arg Ser Trp His Glu Glu Ser Pro Phe Glu Lys Gln Phe Lys
610 615 620

Arg Arg Ser Cys Gln Met Glu Phe Gly Glu Ser Ile Met Ser Glu Asn
625 630 635 640

Arg Ser Arg Glu Glu Leu Gly Lys Val Gly Ser Gln Ser Ser Phe Ser
645 650 655

Gly Ser Met Glu Ile Ile Glu Val Ser
660 665

<210> 248
<211> 301
<212> PRT

<213> Homo sapiens

<400> 248

Met Lys Ser Asn Pro Ala Ile Gln Ala Ala Ile Asp Leu Thr Ala Gly
 1 5 10 15

Ala Ala Gly Gly Thr Ala Cys Val Leu Thr Gly Gln Pro Phe Asp Thr
 20 25 30

Met Lys Val Lys Met Gln Thr Phe Pro Asp Leu Tyr Arg Gly Leu Thr
 35 40 45

Asp Cys Cys Leu Lys Thr Tyr Ser Gln Val Gly Phe Arg Gly Phe Tyr
 50 55 60

Lys Gly Thr Ser Pro Ala Leu Ile Ala Asn Ile Ala Glu Asn Ser Val
 65 70 75 80

Leu Phe Met Cys Tyr Gly Phe Cys Gln Gln Val Val Arg Lys Val Ala
 85 90 95

Gly Leu Asp Lys Gln Ala Lys Leu Ser Asp Leu Gln Asn Ala Ala Ala
 100 105 110

Gly Ser Phe Ala Ser Ala Phe Ala Ala Leu Val Leu Cys Pro Thr Glu
 115 120 125

Leu Val Lys Cys Arg Leu Gln Thr Met Tyr Glu Met Glu Thr Ser Gly
 130 135 140

Lys Ile Ala Lys Ser Gln Asn Thr Val Trp Ser Val Ile Lys Ser Ile
 145 150 155 160

Leu Arg Lys Asp Gly Pro Leu Gly Phe Tyr His Gly Leu Ser Ser Thr
 165 170 175

Leu Leu Arg Glu Val Pro Gly Tyr Phe Phe Phe Phe Gly Gly Tyr Glu
 180 185 190

Leu Ser Arg Ser Phe Phe Ala Ser Gly Arg Ser Lys Asp Glu Leu Gly
 195 200 205

Pro Val Pro Leu Met Leu Ser Gly Gly Val Gly Gly Ile Cys Leu Trp

210 215 220
 Leu Ala Val Tyr Pro Val Asp Cys Ile Lys Ser Arg Ile Gln Val Leu
 225 230 235 240
 Ser Met Ser Gly Lys Gln Ala Gly Phe Ile Arg Thr Phe Ile Asn Val
 245 250 255
 Val Lys Asn Glu Gly Ile Thr Ala Leu Tyr Ser Gly Leu Lys Pro Thr
 260 265 270
 Met Ile Arg Ala Phe Pro Ala Asn Gly Ala Leu Phe Leu Ala Tyr Glu
 275 280 285
 Tyr Ser Arg Lys Leu Met Met Asn Gln Leu Glu Ala Tyr
 290 295 300

 <210> 249
 <211> 337
 <212> PRT
 <213> Homo sapiens

 <400> 249
 Met Ala Ala Pro Arg Asp Asn Val Thr Leu Leu Phe Lys Leu Tyr Cys
 1 5 10 15
 Leu Ala Val Met Thr Leu Met Ala Ala Val Tyr Thr Ile Ala Leu Arg
 20 25 30
 Tyr Thr Arg Thr Ser Asp Lys Glu Leu Tyr Phe Ser Thr Thr Ala Val
 35 40 45
 Cys Ile Thr Glu Val Ile Lys Leu Leu Leu Ser Val Gly Ile Leu Ala
 50 55 60
 Lys Glu Thr Gly Ser Leu Gly Arg Phe Lys Ala Ser Leu Arg Glu Asn
 65 70 75 80
 Val Leu Gly Ser Pro Lys Glu Leu Leu Lys Leu Ser Val Pro Ser Leu
 85 90 95
 Val Tyr Ala Val Gln Asn Asn Met Ala Phe Leu Ala Leu Ser Asn Leu
 100 105 110

Asp Ala Ala Val Tyr Gln Val Thr Tyr Gln Leu Lys Ile Pro Cys Thr
 115 120 125

Ala Leu Cys Thr Val Leu Met Leu Asn Arg Thr Leu Ser Lys Leu Gln
 130 135 140

Trp Val Ser Val Phe Met Leu Cys Ala Gly Val Thr Leu Val Gln Trp
 145 150 155 160

Lys Pro Ala Gln Ala Thr Lys Val Val Val Glu Gln Asn Pro Leu Leu
 165 170 175

Gly Phe Gly Ala Ile Ala Ile Ala Val Leu Cys Ser Gly Phe Ala Gly
 180 185 190

Val Tyr Phe Glu Lys Val Leu Lys Ser Ser Asp Thr Ser Leu Trp Val
 195 200 205

Arg Asn Ile Gln Met Tyr Leu Ser Gly Ile Ile Val Thr Leu Ala Gly
 210 215 220

Val Tyr Leu Ser Asp Gly Ala Glu Ile Lys Glu Lys Gly Phe Phe Tyr
 225 230 235 240

Gly Tyr Thr Tyr Tyr Val Trp Phe Val Ile Phe Leu Ala Ser Val Gly
 245 250 255

Gly Leu Tyr Thr Ser Val Val Val Lys Tyr Thr Asp Asn Ile Met Lys
 260 265 270

Gly Phe Ser Ala Ala Ala Ala Ile Val Leu Ser Thr Ile Ala Ser Val
 275 280 285

Met Leu Phe Gly Leu Gln Ile Thr Leu Thr Phe Ala Leu Gly Thr Leu
 290 295 300

Leu Val Cys Val Ser Ile Tyr Leu Tyr Gly Leu Pro Arg Gln Asp Thr
 305 310 315 320

Thr Ser Ile Gln Gln Gly Glu Thr Ala Ser Lys Glu Arg Val Ile Gly
 325 330 335

Val

<210> 250
 <211> 487
 <212> PRT
 <213> Homo sapiens

<400> 250

Met Met His Phe Lys Ser Gly Leu Glu Leu Thr Glu Leu Gln Asn Met
 1 5 10 15

Thr Val Pro Glu Asp Asp Asn Ile Ser Asn Asp Ser Asn Asp Phe Thr
 20 25 30

Glu Val Glu Asn Gly Gln Ile Asn Ser Lys Phe Ile Ser Asp Arg Glu
 35 40 45

Ser Arg Arg Ser Leu Thr Asn Ser His Leu Glu Lys Lys Lys Cys Asp
 50 55 60

Glu Tyr Ile Pro Gly Thr Thr Ser Leu Gly Met Ser Val Phe Asn Leu
 65 70 75 80

Ser Asn Ala Ile Met Gly Ser Gly Ile Leu Gly Leu Ala Phe Ala Leu
 85 90 95

Ala Asn Thr Gly Ile Leu Leu Phe Leu Val Leu Leu Thr Ser Val Thr
 100 105 110

Leu Leu Ser Ile Tyr Ser Ile Asn Leu Leu Leu Ile Cys Ser Lys Glu
 115 120 125

Thr Gly Cys Met Val Tyr Glu Lys Leu Gly Glu Gln Val Phe Gly Thr
 130 135 140

Thr Gly Lys Phe Val Ile Phe Gly Ala Thr Ser Leu Gln Asn Thr Gly
 145 150 155 160

Ala Met Leu Ser Tyr Leu Phe Ile Val Lys Asn Glu Leu Pro Ser Ala
 165 170 175

Ile Lys Phe Leu Met Gly Lys Glu Glu Thr Phe Ser Ala Trp Tyr Val
 180 185 190

Asp Gly Arg Val Leu Val Val Ile Val Thr Phe Gly Ile Ile Leu Pro
 195 200 205

Leu Cys Leu Leu Lys Asn Leu Gly Tyr Leu Gly Tyr Thr Ser Gly Phe
 210 215 220

Ser Leu Ser Cys Met Val Phe Phe Leu Ile Val Val Ile Tyr Lys Lys
 225 230 235 240

Phe Gln Ile Pro Cys Ile Val Pro Glu Leu Asn Ser Thr Ile Ser Ala
 245 250 255

Asn Ser Thr Asn Ala Asp Thr Cys Thr Pro Lys Tyr Val Thr Phe Asn
 260 265 270

Ser Lys Thr Val Tyr Ala Leu Pro Thr Ile Ala Phe Ala Phe Val Cys
 275 280 285

His Pro Ser Val Leu Pro Ile Tyr Ser Glu Leu Lys Asp Arg Ser Gln
 290 295 300

Lys Lys Met Gln Met Val Ser Asn Ile Ser Phe Phe Ala Met Phe Val
 305 310 315 320

Met Tyr Phe Leu Thr Ala Ile Phe Gly Tyr Leu Thr Phe Tyr Asp Asn
 325 330 335

Val Gln Ser Asp Leu Leu His Lys Tyr Gln Ser Lys Asp Asp Ile Leu
 340 345 350

Ile Leu Thr Val Arg Leu Ala Val Ile Val Ala Val Ile Leu Thr Val
 355 360 365

Pro Val Leu Phe Phe Thr Val Arg Ser Ser Leu Phe Glu Leu Ala Lys
 370 375 380

Lys Thr Lys Phe Asn Leu Cys Arg His Thr Val Val Thr Cys Ile Leu
 385 390 395 400

Leu Val Val Ile Asn Leu Leu Val Ile Phe Ile Pro Ser Met Lys Asp
 405 410 415

Ile Phe Gly Val Val Gly Val Thr Ser Ala Asn Met Leu Ile Phe Ile
 420 425 430

Leu Pro Ser Ser Leu Tyr Leu Lys Ile Thr Asp Gln Asp Gly Asp Lys
 435 440 445

Gly Thr Gln Arg Ile Trp Ala Ala Leu Phe Leu Gly Leu Gly Val Leu
 450 455 460

Phe Ser Leu Val Ser Ile Pro Leu Val Ile Tyr Asp Trp Ala Cys Ser
 465 470 475 480

Ser Ser Ser Asp Glu Gly His
 485

<210> 251
 <211> 528
 <212> PRT
 <213> Homo sapiens

<400> 251

Met Ala Gly Ser Asp Thr Ala Pro Phe Leu Ser Gln Ala Asp Asp Pro
 1 5 10 15

Asp Asp Gly Pro Val Pro Gly Thr Pro Gly Leu Pro Gly Ser Thr Gly
 20 25 30

Asn Pro Lys Ser Glu Glu Pro Glu Val Pro Asp Gln Glu Gly Leu Gln
 35 40 45

Arg Ile Thr Gly Leu Ser Pro Gly Arg Ser Ala Leu Ile Val Ala Val
 50 55 60

Leu Cys Tyr Ile Asn Leu Leu Asn Tyr Met Asp Arg Phe Thr Val Ala
 65 70 75 80

Gly Val Leu Pro Asp Ile Glu Gln Phe Phe Asn Ile Gly Asp Ser Ser
 85 90 95

Ser Gly Leu Ile Gln Thr Val Phe Ile Ser Ser Tyr Met Val Leu Ala
 100 105 110

Pro Val Phe Gly Tyr Leu Gly Asp Arg Tyr Asn Arg Lys Tyr Leu Met

115	120	125
Cys Gly Gly Ile Ala Phe Trp Ser Leu Val Thr Leu Gly Ser Ser Phe		
130	135	140
Ile Pro Gly Glu His Phe Trp Leu Leu Leu Leu Thr Arg Gly Leu Val		
145	150	155 160
Gly Val Gly Glu Ala Ser Tyr Ser Thr Ile Ala Pro Thr Leu Ile Ala		
165	170	175
Asp Leu Phe Val Ala Asp Gln Arg Ser Arg Met Leu Ser Ile Phe Tyr		
180	185	190
Phe Ala Ile Pro Val Gly Ser Gly Leu Gly Tyr Ile Ala Gly Ser Lys		
195	200	205
Val Lys Asp Met Ala Gly Asp Trp His Trp Ala Leu Arg Val Thr Pro		
210	215	220
Gly Leu Gly Val Val Ala Val Leu Leu Leu Phe Leu Val Val Arg Glu		
225	230	235 240
Pro Pro Arg Gly Ala Val Glu Arg His Ser Asp Leu Pro Pro Leu Asn		
245	250	255
Pro Thr Ser Trp Trp Ala Asp Leu Arg Ala Leu Ala Arg Asn Pro Ser		
260	265	270
Phe Val Leu Ser Ser Leu Gly Phe Thr Ala Val Ala Phe Val Thr Gly		
275	280	285
Ser Leu Ala Leu Trp Ala Pro Ala Phe Leu Leu Arg Ser Arg Val Val		
290	295	300
Leu Gly Glu Thr Pro Pro Cys Leu Pro Gly Asp Ser Cys Ser Ser Ser		
305	310	315 320
Asp Ser Leu Ile Phe Gly Leu Ile Thr Cys Leu Thr Gly Val Leu Gly		
325	330	335
Val Gly Leu Gly Val Glu Ile Ser Arg Arg Leu Arg His Ser Asn Pro		
340	345	350

Arg Ala Asp Pro Leu Val Cys Ala Thr Gly Leu Leu Gly Ser Ala Pro
 355 360 365

Phe Leu Phe Leu Ser Leu Ala Cys Ala Arg Gly Ser Ile Val Ala Thr
 370 375 380

Tyr Ile Phe Ile Phe Ile Gly Glu Thr Leu Leu Ser Met Asn Trp Ala
 385 390 395 400

Ile Val Ala Asp Ile Leu Leu Tyr Val Val Ile Pro Thr Arg Arg Ser
 405 410 415

Thr Ala Glu Ala Phe Gln Ile Val Leu Ser His Leu Leu Gly Asp Ala
 420 425 430

Gly Ser Pro Tyr Leu Ile Gly Leu Ile Ser Asp Arg Leu Arg Arg Asn
 435 440 445

Trp Pro Pro Ser Phe Leu Ser Glu Phe Arg Ala Leu Gln Phe Ser Leu
 450 455 460

Met Leu Cys Ala Phe Val Gly Ala Leu Gly Gly Ala Ala Phe Leu Gly
 465 470 475 480

Thr Ala Ile Phe Ile Glu Ala Asp Arg Arg Arg Ala Gln Leu His Val
 485 490 495

Gln Gly Leu Leu His Glu Ala Gly Ser Thr Asp Asp Arg Ile Val Val
 500 505 510

Pro Gln Arg Gly Arg Ser Thr Arg Val Pro Val Ala Ser Val Leu Ile
 515 520 525

<210> 252
 <211> 418
 <212> PRT
 <213> Homo sapiens

<400> 252

Met Ala Pro Thr Gln Gly Pro Arg Ala Pro Leu Glu Phe Gly Gly Pro
 1 5 10 15

Leu Gly Ala Ala Ala Leu Leu Leu Leu Leu Pro Ala Thr Met Phe His
 20 25 30

Leu Leu Leu Ala Ala Arg Ser Gly Pro Ala Arg Leu Leu Gly Pro Pro
 35 40 45

Ala Ser Leu Pro Gly Leu Glu Val Leu Trp Ser Pro Arg Ala Leu Leu
 50 55 60

Leu Trp Leu Ala Trp Leu Gly Leu Gln Ala Ala Leu Tyr Leu Leu Pro
 65 70 75 80

Ala Arg Lys Val Ala Glu Gly Gln Glu Leu Lys Asp Lys Ser Arg Leu
 85 90 95

Arg Tyr Pro Ile Asn Gly Phe Gln Ala Leu Val Leu Thr Ala Leu Leu
 100 105 110

Val Gly Leu Gly Met Ser Ala Gly Leu Pro Leu Gly Ala Leu Pro Glu
 115 120 125

Met Leu Leu Pro Leu Ala Phe Val Ala Thr Leu Thr Ala Phe Ile Phe
 130 135 140

Ser Leu Phe Leu Tyr Met Lys Ala Gln Val Ala Pro Val Ser Ala Leu
 145 150 155 160

Ala Pro Gly Gly Asn Ser Gly Asn Pro Ile Tyr Asp Phe Phe Leu Gly
 165 170 175

Arg Glu Leu Asn Pro Arg Ile Cys Phe Phe Asp Phe Lys Tyr Phe Cys
 180 185 190

Glu Leu Arg Pro Gly Leu Ile Gly Trp Val Leu Ile Asn Leu Ala Leu
 195 200 205

Leu Met Lys Glu Ala Glu Leu Arg Gly Ser Pro Ser Leu Ala Met Trp
 210 215 220

Leu Val Asn Gly Phe Gln Leu Leu Tyr Val Gly Asp Ala Leu Trp His
 225 230 235 240

Glu Glu Ala Val Leu Thr Thr Met Asp Ile Thr His Asp Gly Phe Gly

245 250 255
 Phe Met Leu Ala Phe Gly Asp Met Ala Trp Val Pro Phe Thr Tyr Ser
 260 265 270
 Leu Gln Ala Gln Phe Leu Leu His His Pro Gln Pro Leu Gly Leu Pro
 275 280 285
 Met Ala Ser Val Ile Cys Leu Ile Asn Ala Ile Gly Tyr Tyr Ile Phe
 290 295 300
 Arg Gly Ala Asn Ser Gln Lys Asn Thr Phe Arg Lys Asn Pro Ser Asp
 305 310 315 320
 Pro Arg Val Ala Gly Leu Glu Thr Ile Ser Thr Ala Thr Gly Arg Lys
 325 330 335
 Leu Leu Val Ser Gly Trp Trp Gly Met Val Arg His Pro Asn Tyr Leu
 340 345 350
 Gly Asp Leu Ile Met Ala Leu Ala Trp Ser Leu Pro Cys Gly Val Ser
 355 360 365
 His Leu Leu Pro Tyr Phe Tyr Leu Leu Tyr Phe Thr Ala Leu Leu Val
 370 375 380
 His Arg Glu Ala Arg Asp Glu Arg Gln Cys Leu Gln Lys Tyr Gly Leu
 385 390 395 400
 Ala Trp Gln Glu Tyr Cys Arg Arg Val Pro Tyr Arg Ile Met Pro Tyr
 405 410 415

Ile Tyr

<210> 253
 <211> 1281
 <212> PRT
 <213> Homo sapiens

<400> 253

Met Val Arg Lys Lys Asn Pro Pro Leu Arg Asn Val Ala Ser Glu Gly
 1 5 10 15

Glu Gly Gln Ile Leu Glu Pro Ile Gly Thr Glu Ser Lys Val Ser Gly
 20 25 30

Lys Asn Lys Glu Phe Ser Ala Asp Gln Met Ser Glu Asn Thr Asp Gln
 35 40 45

Ser Asp Ala Ala Glu Leu Asn His Lys Glu Glu His Ser Leu His Val
 50 55 60

Gln Asp Pro Ser Ser Ser Ser Lys Lys Asp Leu Lys Ser Ala Val Leu
 65 70 75 80

Ser Glu Lys Ala Gly Phe Asn Tyr Glu Ser Pro Ser Lys Gly Gly Asn
 85 90 95

Phe Pro Ser Phe Pro His Asp Glu Val Thr Asp Arg Asn Met Leu Ala
 100 105 110

Phe Ser Ser Pro Ala Ala Gly Gly Val Cys Glu Pro Leu Lys Ser Pro
 115 120 125

Gln Arg Ala Glu Ala Asp Asp Pro Gln Asp Met Ala Cys Thr Pro Ser
 130 135 140

Gly Asp Ser Leu Glu Thr Lys Glu Asp Gln Lys Met Ser Pro Lys Ala
 145 150 155 160

Thr Glu Glu Thr Gly Gln Ala Gln Ser Gly Gln Ala Asn Cys Gln Gly
 165 170 175

Leu Ser Pro Val Ser Val Ala Ser Lys Asn Pro Gln Val Pro Ser Asp
 180 185 190

Gly Gly Val Arg Leu Asn Lys Ser Lys Thr Asp Leu Leu Val Asn Asp
 195 200 205

Asn Pro Asp Pro Ala Pro Leu Ser Pro Glu Leu Gln Asp Phe Lys Cys
 210 215 220

Asn Ile Cys Gly Tyr Gly Tyr Tyr Gly Asn Asp Pro Thr Asp Leu Ile
 225 230 235 240

Lys His Phe Arg Lys Tyr His Leu Gly Leu His Asn Arg Thr Arg Gln
 245 250 255

Asp Ala Glu Leu Asp Ser Lys Ile Leu Ala Leu His Asn Met Val Gln
 260 265 270

Phe Ser His Ser Lys Asp Phe Gln Lys Val Asn Arg Ser Val Phe Ser
 275 280 285

Gly Val Leu Gln Asp Ile Asn Ser Ser Arg Pro Val Leu Leu Asn Gly
 290 295 300

Thr Tyr Asp Val Gln Val Thr Ser Gly Gly Thr Phe Ile Gly Ile Gly
 305 310 315 320

Arg Lys Thr Pro Asp Cys Gln Gly Asn Thr Lys Tyr Phe Arg Cys Lys
 325 330 335

Phe Cys Asn Phe Thr Tyr Met Gly Asn Ser Ser Thr Glu Leu Glu Gln
 340 345 350

His Phe Leu Gln Thr His Pro Asn Lys Ile Lys Ala Ser Leu Pro Ser
 355 360 365

Ser Glu Val Ala Lys Pro Ser Glu Lys Asn Ser Asn Lys Ser Ile Pro
 370 375 380

Ala Leu Gln Ser Ser Asp Ser Gly Asp Leu Gly Lys Trp Gln Asp Lys
 385 390 395 400

Ile Thr Val Lys Ala Gly Asp Asp Thr Pro Val Gly Tyr Ser Val Pro
 405 410 415

Ile Lys Pro Leu Asp Ser Ser Arg Gln Asn Gly Thr Glu Ala Thr Ser
 420 425 430

Tyr Tyr Trp Cys Lys Phe Cys Ser Phe Ser Cys Glu Ser Ser Ser Ser
 435 440 445

Leu Lys Leu Leu Glu His Tyr Gly Lys Gln His Gly Ala Val Gln Ser
 450 455 460

Gly Gly Leu Asn Pro Glu Leu Asn Asp Lys Leu Ser Arg Gly Ser Val

465 470 475 480
 Ile Asn Gln Asn Asp Leu Ala Lys Ser Ser Glu Gly Glu Thr Met Thr
 485 490 495
 Lys Thr Asp Lys Ser Ser Ser Gly Ala Lys Lys Lys Asp Phe Ser Ser
 500 505 510
 Lys Gly Ala Glu Asp Asn Met Val Thr Ser Tyr Asn Cys Gln Phe Cys
 515 520 525
 Asp Phe Arg Tyr Ser Lys Ser His Gly Pro Asp Val Ile Val Val Gly
 530 535 540
 Pro Leu Leu Arg His Tyr Gln Gln Leu His Asn Ile His Lys Cys Thr
 545 550 555 560
 Ile Lys His Cys Pro Phe Cys Pro Arg Gly Leu Cys Ser Pro Glu Lys
 565 570 575
 His Leu Gly Glu Ile Thr Tyr Pro Phe Ala Cys Arg Lys Ser Asn Cys
 580 585 590
 Ser His Cys Ala Leu Leu Leu Leu His Leu Ser Pro Gly Ala Ala Gly
 595 600 605
 Ser Ser Arg Val Lys His Gln Cys His Gln Cys Ser Phe Thr Thr Pro
 610 615 620
 Asp Val Asp Val Leu Leu Phe His Tyr Glu Ser Val His Glu Ser Gln
 625 630 635 640
 Ala Ser Asp Val Lys Gln Glu Ala Asn His Leu Gln Gly Ser Asp Gly
 645 650 655
 Gln Gln Ser Val Lys Glu Ser Lys Glu His Ser Cys Thr Lys Cys Asp
 660 665 670
 Phe Ile Thr Gln Val Glu Glu Glu Ile Ser Arg His Tyr Arg Arg Ala
 675 680 685
 His Ser Cys Tyr Lys Cys Arg Gln Cys Ser Phe Thr Ala Ala Asp Thr
 690 695 700

Gln Ser Leu Leu Glu His Phe Asn Thr Val His Cys Gln Glu Gln Asp
705 710 715 720

Ile Thr Thr Ala Asn Gly Glu Glu Asp Gly His Ala Ile Ser Thr Ile
725 730 735

Lys Glu Glu Pro Lys Ile Asp Phe Arg Val Tyr Asn Leu Leu Thr Pro
740 745 750

Asp Ser Lys Met Gly Glu Pro Val Ser Glu Ser Val Val Lys Arg Glu
755 760 765

Lys Leu Glu Glu Lys Asp Gly Leu Lys Glu Lys Val Trp Thr Glu Ser
770 775 780

Ser Ser Asp Asp Leu Arg Asn Val Thr Trp Arg Gly Ala Asp Ile Leu
785 790 795 800

Arg Gly Ser Pro Ser Tyr Thr Gln Ala Ser Leu Gly Leu Leu Thr Pro
805 810 815

Val Ser Gly Thr Gln Glu Gln Thr Lys Thr Leu Arg Asp Ser Pro Asn
820 825 830

Val Glu Ala Ala His Leu Ala Arg Pro Ile Tyr Gly Leu Ala Val Glu
835 840 845

Thr Lys Gly Phe Leu Gln Gly Ala Pro Ala Gly Gly Glu Lys Ser Gly
850 855 860

Ala Leu Pro Gln Gln Tyr Pro Ala Ser Gly Glu Asn Lys Ser Lys Asp
865 870 875 880

Glu Ser Gln Ser Leu Leu Arg Arg Arg Arg Gly Ser Gly Val Phe Cys
885 890 895

Ala Asn Cys Leu Thr Thr Lys Thr Ser Leu Trp Arg Lys Asn Ala Asn
900 905 910

Gly Gly Tyr Val Cys Asn Ala Cys Gly Leu Tyr Gln Lys Leu His Ser
915 920 925

Thr Pro Arg Pro Leu Asn Ile Ile Lys Gln Asn Asn Gly Glu Gln Ile
 930 935 940

Ile Arg Arg Arg Thr Arg Lys Arg Leu Asn Pro Glu Ala Leu Gln Ala
 945 950 955 960

Glu Gln Leu Asn Lys Gln Gln Arg Gly Ser Asn Glu Glu Gln Val Asn
 965 970 975

Gly Ser Pro Leu Glu Arg Arg Ser Glu Asp His Leu Thr Glu Ser His
 980 985 990

Gln Arg Glu Ile Pro Leu Pro Ser Leu Ser Lys Tyr Glu Ala Gln Gly
 995 1000 1005

Ser Leu Thr Lys Ser His Ser Ala Gln Gln Pro Val Leu Val Ser
 1010 1015 1020

Gln Thr Leu Asp Ile His Lys Arg Met Gln Pro Leu His Ile Gln
 1025 1030 1035

Ile Lys Ser Pro Gln Glu Ser Thr Gly Asp Pro Gly Asn Ser Ser
 1040 1045 1050

Ser Val Ser Glu Gly Lys Gly Ser Ser Glu Arg Gly Ser Pro Ile
 1055 1060 1065

Glu Lys Tyr Met Arg Pro Ala Lys His Pro Asn Tyr Ser Pro Pro
 1070 1075 1080

Gly Ser Pro Ile Glu Lys Tyr Gln Tyr Pro Leu Phe Gly Leu Pro
 1085 1090 1095

Phe Val His Asn Asp Phe Gln Ser Glu Ala Asp Trp Leu Arg Phe
 1100 1105 1110

Trp Ser Lys Tyr Lys Leu Ser Val Pro Gly Asn Pro His Tyr Leu
 1115 1120 1125

Ser His Val Pro Gly Leu Pro Asn Pro Cys Gln Asn Tyr Val Pro
 1130 1135 1140

Tyr Pro Thr Phe Asn Leu Pro Pro His Phe Ser Ala Val Gly Ser
1145 1150 1155

Asp Asn Asp Ile Pro Leu Asp Leu Ala Ile Lys His Ser Arg Pro
1160 1165 1170

Gly Pro Thr Ala Asn Gly Ala Ser Lys Glu Lys Thr Lys Ala Pro
1175 1180 1185

Pro Asn Val Lys Asn Glu Gly Pro Leu Asn Val Val Lys Thr Glu
1190 1195 1200

Lys Val Asp Arg Ser Thr Gln Asp Glu Leu Ser Thr Lys Cys Val
1205 1210 1215

His Cys Gly Ile Val Phe Leu Asp Glu Val Met Tyr Ala Leu His
1220 1225 1230

Met Ser Cys His Gly Asp Ser Gly Pro Phe Gln Cys Ser Ile Cys
1235 1240 1245

Gln His Leu Cys Thr Asp Lys Tyr Asp Phe Thr Thr His Ile Gln
1250 1255 1260

Arg Gly Leu His Arg Asn Asn Ala Gln Val Glu Lys Asn Gly Lys
1265 1270 1275

Pro Lys Glu
1280

<210> 254

<211> 822

<212> PRT

<213> Homo sapiens

<400> 254

Met Val Ser Trp Gly Arg Phe Ile Cys Leu Val Val Val Thr Met Ala
1 5 10 15

Thr Leu Ser Leu Ala Arg Pro Ser Phe Ser Leu Val Glu Asp Thr Thr
20 25 30

Leu Glu Pro Glu Glu Pro Pro Thr Lys Tyr Gln Ile Ser Gln Pro Glu
35 40 45

Val Tyr Val Ala Ala Pro Gly Glu Ser Leu Glu Val Arg Cys Leu Leu
50 55 60

Lys Asp Ala Ala Val Ile Ser Trp Thr Lys Asp Gly Val His Leu Gly
65 70 75 80

Pro Asn Asn Arg Thr Val Leu Ile Gly Glu Tyr Leu Gln Ile Lys Gly
85 90 95

Ala Thr Pro Arg Asp Ser Gly Leu Tyr Ala Cys Thr Ala Ser Arg Thr
100 105 110

Val Asp Ser Glu Thr Trp Tyr Phe Met Val Asn Val Thr Asp Ala Ile
115 120 125

Ser Ser Gly Asp Asp Glu Asp Asp Thr Asp Gly Ala Glu Asp Phe Val
130 135 140

Ser Glu Asn Ser Asn Asn Lys Arg Ala Pro Tyr Trp Thr Asn Thr Glu
145 150 155 160

Lys Met Glu Lys Arg Leu His Ala Val Pro Ala Ala Asn Thr Val Lys
165 170 175

Phe Arg Cys Pro Ala Gly Gly Asn Pro Met Pro Thr Met Arg Trp Leu
180 185 190

Lys Asn Gly Lys Glu Phe Lys Gln Glu His Arg Ile Gly Gly Tyr Lys
195 200 205

Val Arg Asn Gln His Trp Ser Leu Ile Met Glu Ser Val Val Pro Ser
210 215 220

Asp Lys Gly Asn Tyr Thr Cys Val Val Glu Asn Glu Tyr Gly Ser Ile
225 230 235 240

Asn His Thr Tyr His Leu Asp Val Val Glu Arg Ser Pro His Arg Pro
245 250 255

Ile Leu Gln Ala Gly Leu Pro Ala Asn Ala Ser Thr Val Val Gly Gly
260 265 270

Asp Val Glu Phe Val Cys Lys Val Tyr Ser Asp Ala Gln Pro His Ile
 275 280 285

Gln Trp Ile Lys His Val Glu Lys Asn Gly Ser Lys Tyr Gly Pro Asp
 290 295 300

Gly Leu Pro Tyr Leu Lys Val Leu Lys His Ser Gly Ile Asn Ser Ser
 305 310 315 320

Asn Ala Glu Val Leu Ala Leu Phe Asn Val Thr Glu Ala Asp Ala Gly
 325 330 335

Glu Tyr Ile Cys Lys Val Ser Asn Tyr Ile Gly Gln Ala Asn Gln Ser
 340 345 350

Ala Trp Leu Thr Val Leu Pro Lys Gln Gln Ala Pro Gly Arg Glu Lys
 355 360 365

Glu Ile Thr Ala Ser Pro Asp Tyr Leu Glu Ile Ala Ile Tyr Cys Ile
 370 375 380

Gly Val Phe Leu Ile Ala Cys Met Val Val Thr Val Ile Leu Cys Arg
 385 390 395 400

Met Lys Asn Thr Thr Lys Lys Pro Asp Phe Ser Ser Gln Pro Ala Val
 405 410 415

His Lys Leu Thr Lys Arg Ile Pro Leu Arg Arg Gln Val Thr Val Ser
 420 425 430

Ala Glu Ser Ser Ser Ser Met Asn Ser Asn Thr Pro Leu Val Arg Ile
 435 440 445

Thr Thr Arg Leu Ser Ser Thr Ala Asp Thr Pro Met Leu Ala Gly Val
 450 455 460

Ser Glu Tyr Glu Leu Pro Glu Asp Pro Lys Trp Glu Phe Pro Arg Asp
 465 470 475 480

Lys Leu Thr Leu Gly Lys Pro Leu Gly Glu Gly Cys Phe Gly Gln Val
 485 490 495

Val Met Ala Glu Ala Val Gly Ile Asp Lys Asp Lys Pro Lys Glu Ala
 500 505 510

Val Thr Val Ala Val Lys Met Leu Lys Asp Asp Ala Thr Glu Lys Asp
 515 520 525

Leu Ser Asp Leu Val Ser Glu Met Glu Met Met Lys Met Ile Gly Lys
 530 535 540

His Lys Asn Ile Ile Asn Leu Leu Gly Ala Cys Thr Gln Asp Gly Pro
 545 550 555 560

Leu Tyr Val Ile Val Glu Tyr Ala Ser Lys Gly Asn Leu Arg Glu Tyr
 565 570 575

Leu Arg Ala Arg Arg Pro Pro Gly Met Glu Tyr Ser Tyr Asp Ile Asn
 580 585 590

Arg Val Pro Glu Glu Gln Met Thr Phe Lys Asp Leu Val Ser Cys Thr
 595 600 605

Tyr Gln Leu Ala Arg Gly Met Glu Tyr Leu Ala Ser Gln Lys Cys Ile
 610 615 620

His Arg Asp Leu Ala Ala Arg Asn Val Leu Val Thr Glu Asn Asn Val
 625 630 635 640

Met Lys Ile Ala Asp Phe Gly Leu Ala Arg Asp Ile Asn Asn Ile Asp
 645 650 655

Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu Pro Val Lys Trp Met Ala
 660 665 670

Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr His Gln Ser Asp Val Trp
 675 680 685

Ser Phe Gly Val Leu Met Trp Glu Ile Phe Thr Leu Gly Gly Ser Pro
 690 695 700

Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe Lys Leu Leu Lys Glu Gly
 705 710 715 720

His Arg Met Asp Lys Pro Ala Asn Cys Thr Asn Glu Leu Tyr Met Met

725 730 735
 Met Arg Asp Cys Trp His Ala Val Pro Ser Gln Arg Pro Thr Phe Lys
 740 745 750
 Gln Leu Val Glu Asp Leu Asp Arg Ile Leu Thr Leu Thr Thr Asn Glu
 755 760 765
 Glu Tyr Leu Asp Leu Ser Gln Pro Leu Glu Gln Tyr Ser Pro Ser Tyr
 770 775 780
 Pro Asp Thr Arg Ser Ser Cys Ser Ser Gly Asp Asp Ser Val Phe Ser
 785 790 795 800
 Pro Asp Pro Met Pro Tyr Glu Pro Cys Leu Pro Gln Tyr Pro His Ile
 805 810 815
 Asn Gly Ser Val Lys Thr
 820
 <210> 255
 <211> 167
 <212> PRT
 <213> Homo sapiens
 <400> 255
 Met Leu Val Leu Leu Ala Phe Ile Ile Ala Phe His Ile Thr Ser Ala
 1 5 10 15
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 Ile Phe Val Leu Gln Leu Phe Arg Leu Lys Gln Gly Glu Arg Phe Val
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Ala Ser Ile Tyr Thr Asp Arg Arg Glu Asp Ile His Asp Lys Asn Ala
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